

ELECTROLUMINESCENT DEVICE

The present invention relates to organo-electroluminescent (EL) devices, in particular EL devices that comprise durable, blue-emitting organo-electroluminescent layers. The organo-
5 electroluminescent layers comprise certain organic compounds containing one or more pyrimidine moieties.

Progress has been made towards developing organic-based electroluminescent devices suitable for full color displays. Generally, an EL device is comprised of a light-emitting layer
10 or layers and a pair of facing electrodes sandwiching the light-emitting layer(s). Application of an electric field between the electrodes results in the injection of electrons and holes to the system, resulting in the release of energy as light.

However, organo EL devices have not been developed that have suitable stability under
15 continuous operation. In particular, there remains a need for blue-emitting, stable organo EL devices.

U.S. Pat. No. 5,104,740 teaches an electroluminescent element that comprises a fluorescent layer containing a coumarinic or azacoumarinic derivative and a hole transport layer, both
20 made of organic compounds and laminated on top of the other.

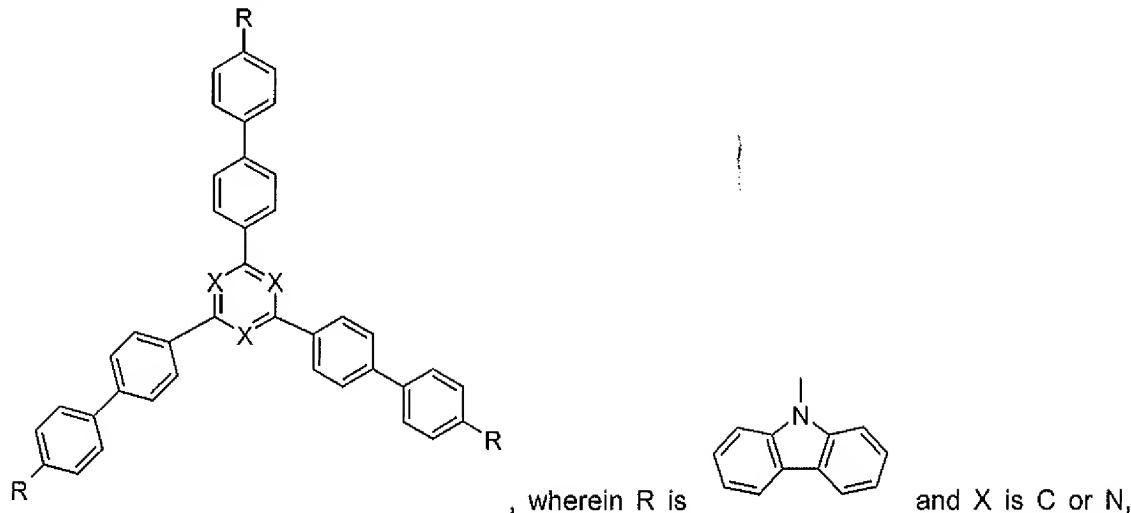
U.S. Pat. No. 6,280,859 discloses certain polyaromatic organic compounds for use as a light-emitting material in organo-electroluminescent devices.

25 U.S. Pat. No. 5,116,708 is aimed at a hole transport material for EL devices.

WO98/04007 and EP-A-1013740 relate to an electroluminescent arrangement with the electron-conducting layer containing one or more compounds comprising triazine as basic substance.

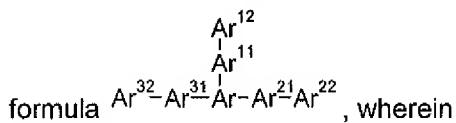
30 EP-A-1013740 discloses the use of triazine compounds in EL devices.

EP-A-1,202,608 discloses EL devices comprising a carbazole compound of formula



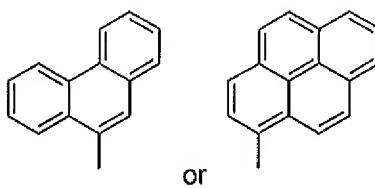
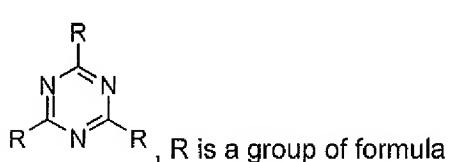
which constitutes the hole transporting layer.

5 JP2002324678 relates to light emitting elements comprising at least one kind of compound of



Ar^{11} , Ar^{21} and Ar^{31} denote arylene groups, Ar^{12} , Ar^{22} and Ar^{32} denote substituents or hydrogen atoms, wherein at least one of Ar^{11} , Ar^{21} , Ar^{31} , Ar^{12} , Ar^{22} and Ar^{32} is either a condensed ring aryl structure or a condensed ring heteroaryl structure; Ar denotes an arylene group or a

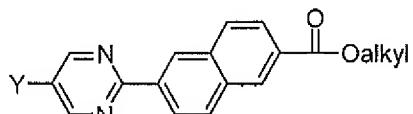
10 heteroarylene group; and at least one amine derivative having a condensed ring group with two or more rings are contained in a luminous layer. As examples of compounds of the above formula, wherein Ar denotes a heteroarylene group the following two compounds are explicitly mentioned:



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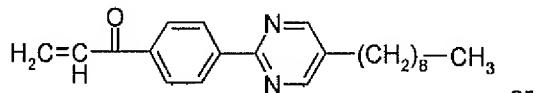
WO02/02714 relates to electroluminescent iridium compounds with fluorinated phenylpyridines, phenylpyrimidines, and phenylquinolines and devices made with such compounds.

US-A-5,770,108 describes liquid crystal compositions comprising pyrimidine compounds of

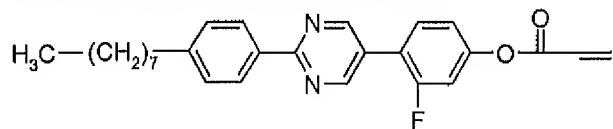


the following formula, wherein Y is alkyl or -O-alkyl and liquid crystal element comprising said composition.

5 WO01/05863 relates to EL devices comprising arylamine-substituted poly(arylene vinylenes).

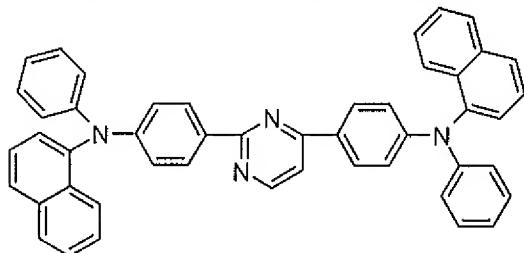


JP2000347432 describes the use of



in EL devices.

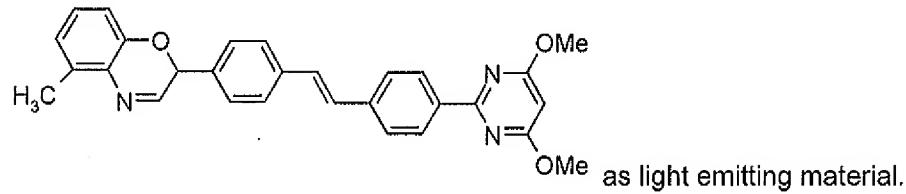
10 EP-A-926216 relates to EL devices using triaryl amine compounds, such as



EP-A-690 053 relates to the use of conjugated compounds containing two or more pyrimidine rings, which are part of the conjugated system, as electroluminescent materials. The

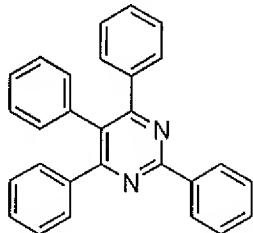
15 conjugated compounds described in EP-A-690 053 comprise pyrimidin-2,5-diyl groups which do not carry substituents at positions 4 and 6.

EP-A-563009 relates to EL devices comprising



as light emitting material.

US-A-5,077,142 relates to EL devices comprising a number of organic compounds as light



emitting material. A pyrimidine moiety, , is listed among a long list of possible organic compounds.

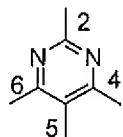
5 It is the object of the present invention to provide a light emitting element with excellent light emitting characteristics and durability.

Certain organic compounds containing one or more pyrimidine moieties are found to be suitable for use in organo-electroluminescent devices. In particular, certain pyrimidine

10 derivatives are suitable blue emitters with good durability.

The present invention is aimed at an electroluminescent device comprising an organic light-emitting layer that contains at least one blue-emitting organic compound containing one or more pyrimidine moieties.

15 Accordingly the present invention relates to an electroluminescent device comprising an anode, a cathode and one or a plurality of organic compound layers sandwiched therebetween, in which said organic compound layers comprise an organic compound containing one or more pyrimidine moieties:



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For example, the present organic compounds comprise one, two, three or more pyrimidine moieties, including oligomers. It is understood that the open valences in the pyrimidine moiety represents a covalent bond that is not limited in its substitution.

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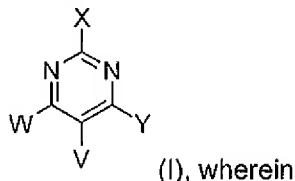
In general, the organic compound or compounds emit light below about 520 nm, in particular between about 380 nm and about 520 nm.

The organic compound or compounds have especially a NTSC coordinate of between about (0.12, 0.05) and about (0.16, 0.10), more especially a NTSC coordinate of about (0.14, 0.08).

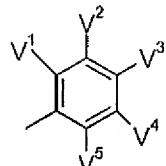
The organic compound or compounds have a melting point above about 150°C, preferably

5 above about 200°C and most preferred above about 250°C.

For example, the organic compound is a pyrimidine compound of formula

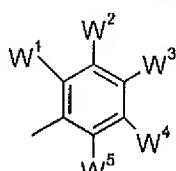


V is C₆-C₃₀aryl or C₂-C₃₀heteroaryl, which can be substituted or unsubstituted, in particular



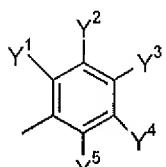
10 , H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl, C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy; C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; -NR⁵R⁶;

W is C₆-C₃₀aryl or C₂-C₃₀heteroaryl, which can be substituted or unsubstituted, in particular



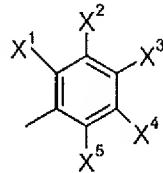
15 , H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl, C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy; C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; -NR⁵R⁶;

Y is C₆-C₃₀aryl or C₂-C₃₀heteroaryl, which can be substituted or unsubstituted, in particular



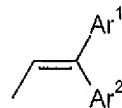
20 , H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl, C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy; C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; -NR⁵R⁶;

X is C₆-C₃₀aryl or C₂-C₃₀heteroaryl, which can be substituted or unsubstituted, in particular



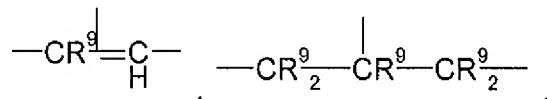
, H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl, C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy; C₁-C₁₈alkoxy

5 which is substituted by E and/or interrupted by D; -SR⁶; -NR⁵R⁶; wherein the groups V¹ to V⁵, W¹ to W⁵, X¹ to X⁵ and Y¹ to Y⁵ are independently of each other H; halogen, C₆-C₂₄aryl; C₆-C₂₄aryl which is substituted by G; C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₇-C₁₈alkylaryl; C₇-C₁₈alkylaryl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl; C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D;

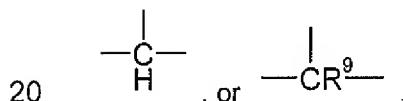


10 , wherein Ar¹ is C₆-C₃₀aryl or C₂-C₃₀heteroaryl, especially phenyl, Ar² is C₆-C₃₀aryl or C₂-C₃₀heteroaryl, especially phenyl, or H, C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy, C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; -NR⁵R⁶; C₂-C₂₄hetero-aryl; C₂-C₂₄heteroaryl which is substituted by L; -SOR⁴; -SO₂R⁴; -COR⁸; -COOR⁷; -CONR⁵R⁶; C₄-C₁₈cycloalkyl; C₄-C₁₈cycloalkyl which is substituted by E and/or interrupted by D; C₄-C₁₈cycloalkenyl; C₄-C₁₈cycloalkenyl which is substituted by E and/or interrupted by D; or

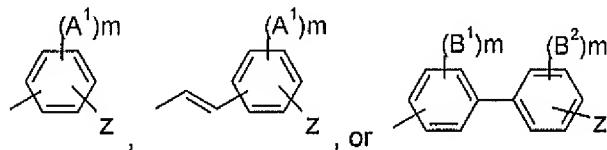
15 W⁵ or Y⁵ together with V form a group -CR⁹₂-, -CR⁹₂-CR⁹₂-, -C(=O)CR⁹₂-, -C(=O)-, or -CR⁹=CR⁹-, or



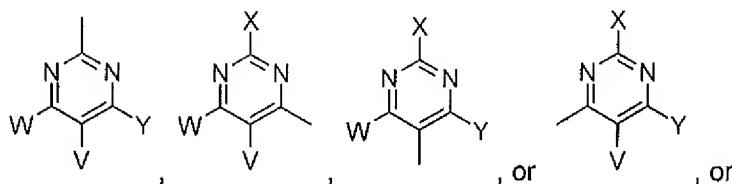
W⁵ and Y⁵ together with V form a group



wherein R⁹ is H; C₁-C₁₈alkyl, C₁-C₁₈alkyl which is interrupted by -O-, C₆-C₁₈aryl, C₆-C₁₈aryl which is substituted by C₁-C₁₈alkyl, or C₁-C₁₈alkoxy, or one of the substituents V, W, X, or Y is a group of the formula -Z, -Ar-Z, wherein Ar is C₆-C₂₄aryl or C₂-C₂₄heteroaryl, which can be substituted, in particular

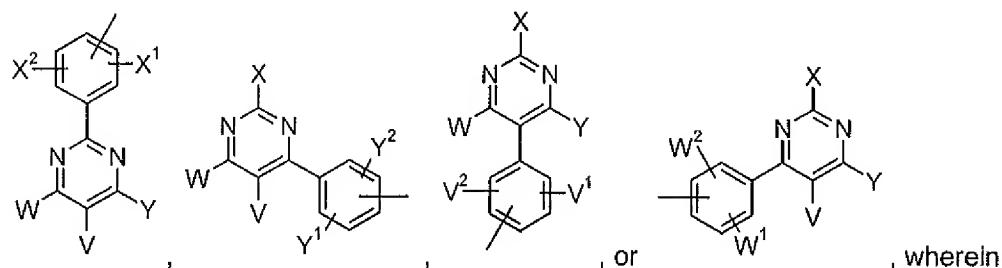
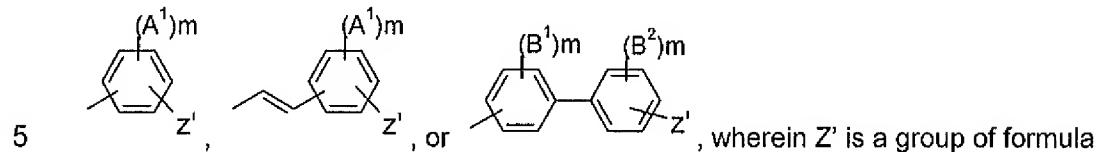


25 , wherein Z is a group of formula



one of the substituents

V^1 to V^5 , W^1 to W^5 , X^1 to X^5 , or Y^1 to Y^5 is a group of the formula $-Z'$, $-Ar-Z'$, wherein Ar is C_6 - C_{24} aryl or C_2 - C_{24} heteroaryl, which can be substituted, in particular



A^1 , B^1 and B^2 are independently of each other H; C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by G; C_1 - C_{18} alkyl; C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D; C_7 - C_{18} alkylaryl; C_7 - C_{18} alkylaryl which is substituted by E and/or interrupted by D; C_2 - C_{18} alkenyl;

10 C_2 - C_{18} alkenyl which is substituted by E and/or interrupted by D; C_2 - C_{18} alkynyl; C_2 - C_{18} alkynyl which is substituted by E and/or interrupted by D; C_1 - C_{18} alkoxy, C_1 - C_{18} alkoxy which is substituted by E and/or interrupted by D; $-SR^5$; $-NR^5R^6$; C_2 - C_{18} heteroaryl; C_2 - C_{18} heteroaryl which is substituted by L; $-SOR^4$; $-SO_2R^4$; $-COR^8$; $-COOR^7$; $-CONR^5R^6$; C_4 - C_{18} cycloalkyl; C_4 - C_{18} cycloalkyl which is substituted by E and/or interrupted by D; C_4 - C_{18} cycloalkenyl; C_4 -

15 C_{18} cycloalkenyl which is substituted by E and/or interrupted by D; or

two substituents A^1 , B^1 , B^2 or B^1 and B^2 form a five to seven membered ring, which can be substituted,

m is an integer of 1 to 4; and W^1 , W^2 , Y^1 , Y^2 , X^1 , X^2 , V , W , X and Y are as defined above;

D is $-CO$; $-COO$; $-OCOO$; $-S$; $-SO$; $-SO_2$; $-O$; $-NR^5$; $-SiR^5R^6$; $-POR^5$; $-CR^5=CR^6$; or -

20 $C\equiv C$;

E is $-OR^5$; $-SR^5$; $-NR^5R^6$; $-COR^8$; $-COOR^7$; $-CONR^5R^6$; $-CN$; $-OCOOR^7$; or halogen, especially F;

G is E; K; heteroaryl; heteroaryl which is substituted by C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by E and/or K;

25 K is C_1 - C_{18} alkyl; C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D; C_7 - C_{18} alkylaryl which is substituted by E and/or interrupted by D; C_2 - C_{18} alkenyl; C_2 - C_{18} alkenyl which is

substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy, C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; C₄-C₁₈cycloalkyl; C₄-C₁₈cycloalkyl which is substituted by E and/or interrupted by D; C₄-C₁₈cycloalkenyl; or C₄-C₁₈cycloalkenyl which is substituted by E and/or interrupted by D;

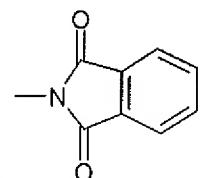
5 interrupted by D;

L is E; K; C₆-C₁₈aryl; or C₆-C₁₈aryl which is substituted by G, E and/or K;

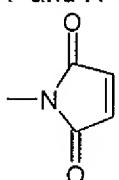
R⁴ is C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by C₁-C₁₈alkyl, C₁-C₁₈alkoxy; C₁-C₁₈alkyl; or C₁-C₁₈alkyl which is interrupted by -O-;

R⁵ and R⁶ are independently of each other H; C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by

10 C₁-C₁₈alkyl, C₁-C₁₈alkoxy; C₁-C₁₈alkyl; or C₁-C₁₈alkyl which is interrupted by -O-; or



or

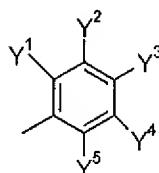


R⁷ is H; C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by C₁-C₁₈alkyl, C₁-C₁₈alkoxy; C₁-C₁₈alkyl; C₁-C₁₈alkyl which is interrupted by -O-;

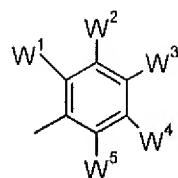
15 R⁸ is H; C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by C₁-C₁₈alkyl, C₁-C₁₈alkoxy; C₁-C₁₈alkyl; C₁-C₁₈alkyl which is interrupted by -O-.

or two substituents selected from V¹ to V⁵, W¹ to W⁵, X¹ to X⁵, Y¹ to Y⁵ which are in neighborhood to each other form a five to seven membered ring.

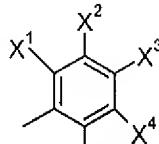
20 Preferably at least one, more preferably at least two, most preferably at least three of the groups V, W, X and Y are C₆-C₃₀aryl or C₂-C₃₀heteroaryl, which optionally can be substituted.



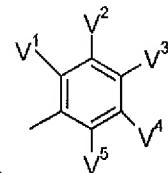
Preferred are compounds of formula I, wherein Y is , wherein Y¹ to Y⁵ are as defined above and at least one of the substituents Y¹ to Y⁵ is different from H; W is



, wherein W^1 to W^5 are as defined above and at least one of the substituents

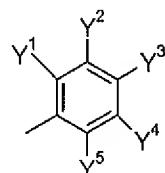


W^1 to W^5 is different from H; X is , wherein X^1 to X^5 are as defined above and at

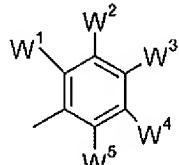


least one of the substituents X^1 to X^5 is different from H, and/or V is , wherein V^1 to V^5 are as defined above and at least one substituent V^1 to V^5 is different from H.

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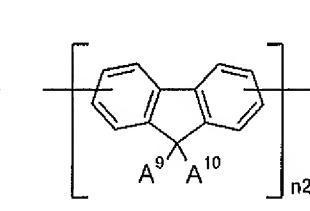
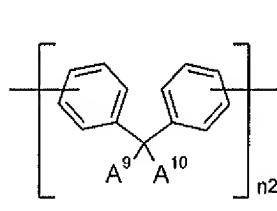
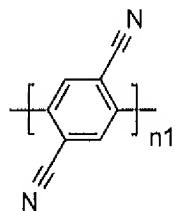
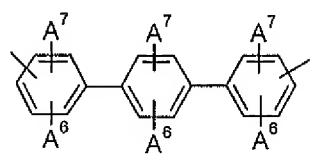
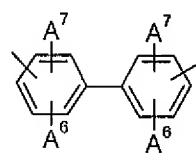
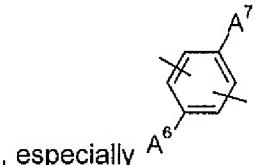
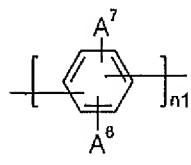


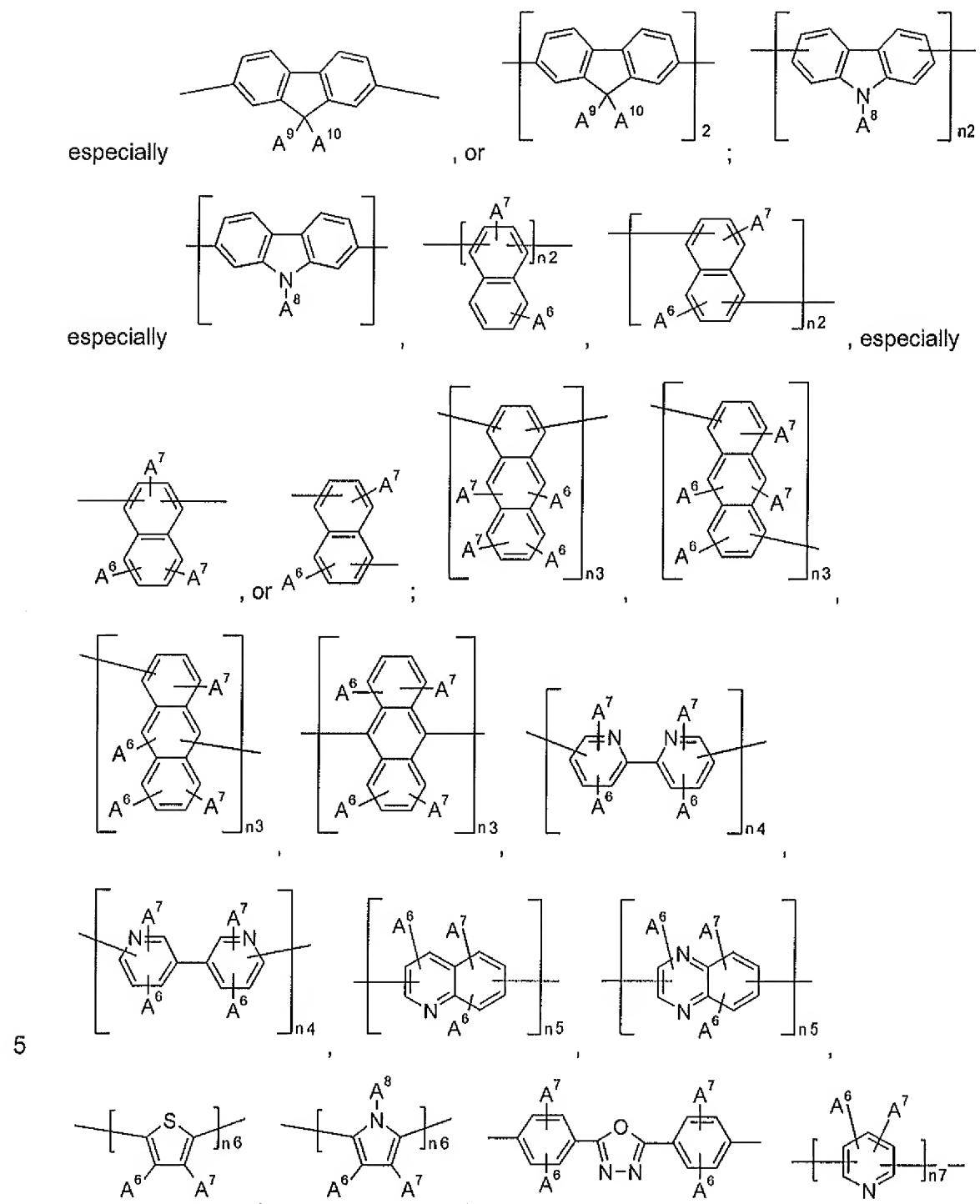
In one further embodiment compounds of formula I are preferred, wherein Y is

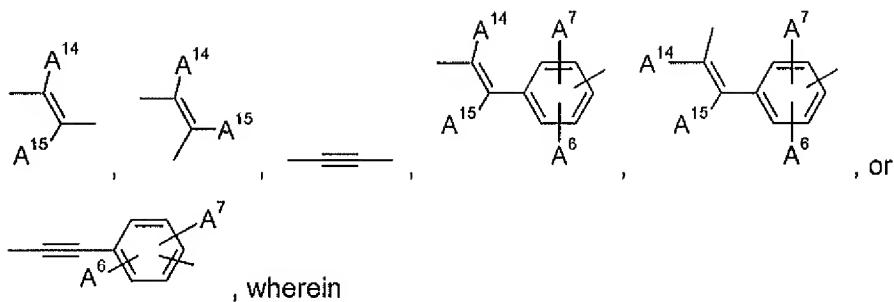


and W is , wherein W^1 and W^5 and Y^1 and Y^5 are independently of each other H; C₁-C₁₈alkyl; or C₁-C₁₈alkyl which is substituted by E and/or interrupted by D.

10 Examples of suitable groups Ar are







n1, n2, n3, n4, n5, n6 and n7 are integers of 1 to 10, in particular 1 to 3,

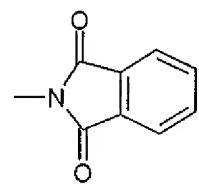
A⁶ and A⁷ are independently of each other H, C₁-C₁₈alkyl, C₁-C₁₈alkyl which is substituted by

5 E and/or interrupted by D, C₆-C₂₄aryl, C₆-C₂₄aryl which is substituted by E, C₂-C₂₀heteroaryl, C₂-C₂₀heteroaryl which is substituted by E, C₂-C₁₈alkenyl, C₂-C₁₈alkynyl, C₁-C₁₈alkoxy, C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D, C₇-C₂₅aralkyl, or -CO-R²⁸, A⁸ is C₁-C₁₈alkyl, C₁-C₁₈alkyl which is substituted by E and/or interrupted by D, C₆-C₂₄aryl, or C₇-C₂₅aralkyl,

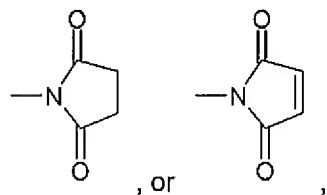
10 A⁹ and A¹⁰ are independently of each other C₁-C₁₈alkyl, C₁-C₁₈alkyl which is substituted by E and/or interrupted by D, C₆-C₂₄aryl, C₆-C₂₄aryl which is substituted by E, C₂-C₂₀heteroaryl, C₂-C₂₀heteroaryl which is substituted by E, C₂-C₁₈alkenyl, C₂-C₁₈alkynyl, C₁-C₁₈alkoxy, C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D, or C₇-C₂₅aralkyl, or A⁹ and A¹⁰ form a ring, especially a five- or six-membered ring,

15 A¹⁴ and A¹⁵ are independently of each other H, C₁-C₁₈alkyl, C₁-C₁₈alkyl which is substituted by E and/or interrupted by D, C₆-C₂₄aryl, C₆-C₂₄aryl which is substituted by E, C₂-C₂₀heteroaryl, or C₂-C₂₀heteroaryl which is substituted by E, D is -CO-; -COO-; -S-; -SO-; -SO₂-; -O-; -NA²⁵-; -SiA³⁰A³¹-; -POA³²-; -CA²³=CA²⁴-; or -C≡C-; and

20 E is -OA²⁹; -SA²⁹; -NA²⁵A²⁶; -COA²⁸; -COOA²⁷; -CONA²⁵A²⁶; -CN; -OCOOA²⁷; or halogen; wherein A²³, A²⁴, A²⁵ and A²⁶ are independently of each other H; C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by C₁-C₁₈alkyl, C₁-C₁₈alkoxy; C₁-C₁₈alkyl; or C₁-C₁₈alkyl which is interrupted by



-O-; or A²⁵ and A²⁶ together form a five or six membered ring, in particular



A^{27} and A^{28} are independently of each other H; C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by C_1 - C_{18} alkyl, or C_1 - C_{18} alkoxy; C_1 - C_{18} alkyl; or C_1 - C_{18} alkyl which is interrupted by $-O-$, A^{29} is H; C_6 - C_{18} aryl; C_6 - C_{18} aryl, which is substituted by C_1 - C_{18} alkyl, C_1 - C_{18} alkoxy; C_1 - C_{18} alkyl; or C_1 - C_{18} alkyl which is interrupted by $-O-$,

5 A^{30} and A^{31} are independently of each other C_1 - C_{18} alkyl, C_6 - C_{18} aryl, or C_6 - C_{18} aryl, which is substituted by C_1 - C_{18} alkyl, and A^{32} is C_1 - C_{18} alkyl, C_6 - C_{18} aryl, or C_6 - C_{18} aryl, which is substituted by C_1 - C_{18} alkyl. Preferably, A^6 and A^7 are independently of each other H, C_1 - C_{18} alkyl, such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, isobutyl, sec-butyl, t-butyl, 2-methylbutyl, n-pentyl, isopentyl, n-hexyl, 2-ethylhexyl, or n-heptyl, C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D, such as $-CH_2OCH_3$, $-CH_2OCH_2CH_3$, $-CH_2OCH_2CH_2OCH_3$, or $-CH_2OCH_2CH_2OCH_2CH_3$, C_6 - C_{24} aryl, such as phenyl, naphthyl, or biphenyl, C_6 - C_{24} aryl which is substituted by E, such as $-C_6H_4OCH_3$, $-C_6H_4OCH_2CH_3$, $-C_6H_3(OCH_3)_2$, or $-C_6H_3(OCH_2CH_3)_2$, $-C_6H_4CH_3$, $-C_6H_3(CH_3)_2$, $-C_6H_2(CH_3)_3$, or $-C_6H_4tBu$.

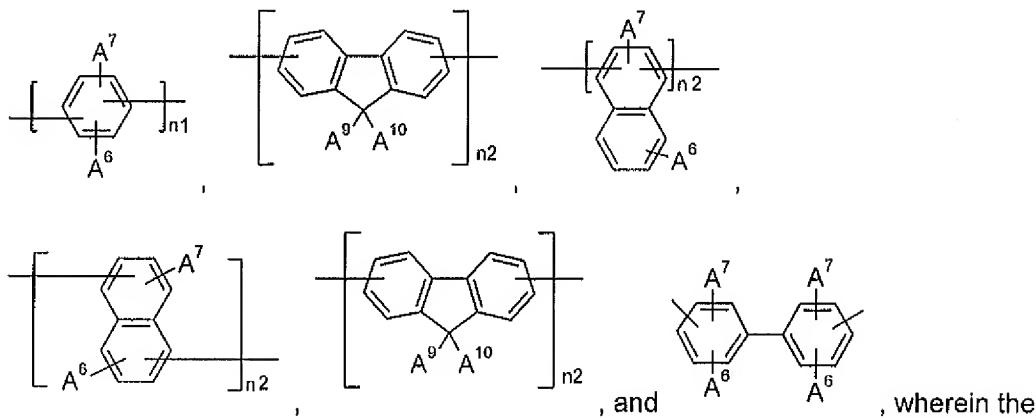
10 A^8 is preferably H, C_1 - C_{18} alkyl, such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, isobutyl, sec-butyl, t-butyl, 2-methylbutyl, n-pentyl, isopentyl, n-hexyl, 2-ethylhexyl, n-heptyl, or C_6 - C_{24} aryl, such as phenyl, naphthyl, or biphenyl. Preferably, A^9 and A^{10} are independently of each other H, C_1 - C_{18} alkyl, such as n-butyl, sec-butyl, hexyl, octyl, or 2-ethyl-hexyl, C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D, such as $-CH_2(OCH_2CH_2)_wOCH_3$, $w = 1, 2, 3$, or 4, C_6 - C_{24} aryl, such as phenyl, naphthyl, or biphenyl, C_6 - C_{24} aryl which is substituted by E, such as $-C_6H_4OCH_3$, $-C_6H_4OCH_2CH_3$, $-C_6H_3(OCH_3)_2$, $-C_6H_3(OCH_2CH_3)_2$, $-C_6H_4CH_3$, $-C_6H_3(CH_3)_2$, $-C_6H_2(CH_3)_3$, or $-C_6H_4tBu$, or A^9 and A^{10} together form a 4 to 8 membered ring, especially a 5 or 6 membered ring, such as cyclohexyl, or cyclopentyl.

15 A^{14} and A^{15} are independently of each other H, C_1 - C_{18} alkyl, such as as methyl, ethyl, n-propyl, iso-propyl, n-butyl, isobutyl, or sec-butyl, or C_6 - C_{24} aryl, such as phenyl, naphthyl, or biphenyl. D is preferably $-CO-$, $-COO-$, $-S-$, $-SO-$, $-SO_2-$, $-O-$, $-NA^{25}-$, wherein A^{25} is C_1 - C_{18} alkyl, such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, isobutyl, or sec-butyl, or C_6 - C_{24} aryl, such as phenyl, naphthyl, or biphenyl.

20 A^{26} is preferably $-OA^{29}$; $-SA^{29}$; $-NA^{25}A^{25}$; $-COA^{28}$; $-COOA^{27}$; $-CONA^{25}A^{25}$; or $-CN$; wherein A^{25} , A^{27} , A^{28} and A^{29} are independently of each other C_1 - C_{18} alkyl, such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, isobutyl, sec-butyl, hexyl, octyl, or 2-ethyl-hexyl, or C_6 - C_{24} aryl, such as phenyl, naphthyl, or biphenyl.

25 E is preferably $-OA^{29}$; $-SA^{29}$; $-NA^{25}A^{25}$; $-COA^{28}$; $-COOA^{27}$; $-CONA^{25}A^{25}$; or $-CN$; wherein A^{25} , A^{27} , A^{28} and A^{29} are independently of each other C_1 - C_{18} alkyl, such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, isobutyl, sec-butyl, hexyl, octyl, or 2-ethyl-hexyl, or C_6 - C_{24} aryl, such as phenyl, naphthyl, or biphenyl.

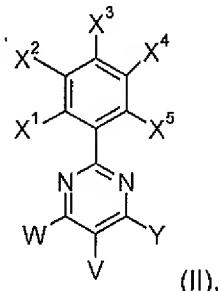
Among the above-mentioned Ar the following groups are preferred:



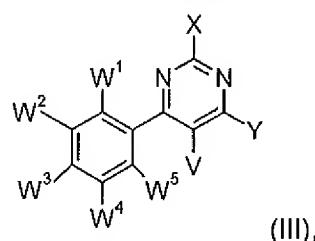
substituents are defined as above.

5

Preferably the electroluminescent device comprises pyrimidine compounds of formula II to VI:



wherein V, W, Y and X¹ to X⁵ are as defined above;



10

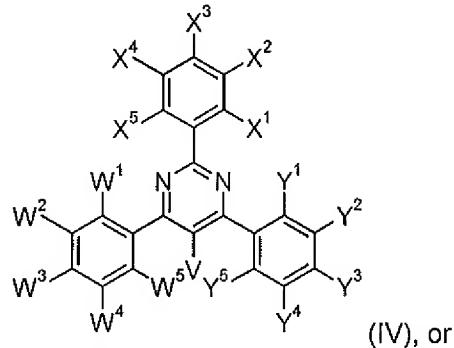
wherein V, X, Y and W¹ to W⁵ are as defined above, especially



Y is R¹, if X is , or X is R¹, if Y is , wherein R¹ is H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl, C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted

15 by E and/or interrupted by D; C₁-C₁₈alkoxy; C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁶; or -NR⁵R⁶; wherein W¹ to W⁵, X¹ to X⁵, Y¹ to Y⁵, E, D, R⁵ and R⁶ are

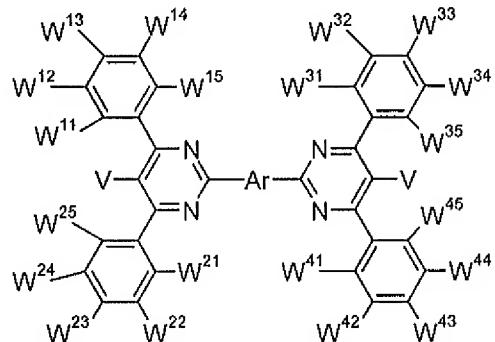
as defined above, wherein most preferred W¹ and W⁵ and Y¹ and Y⁵ are independently of each other H; C₁-C₁₈alkyl; or C₁-C₁₈alkyl which is substituted by E and/or interrupted by D;



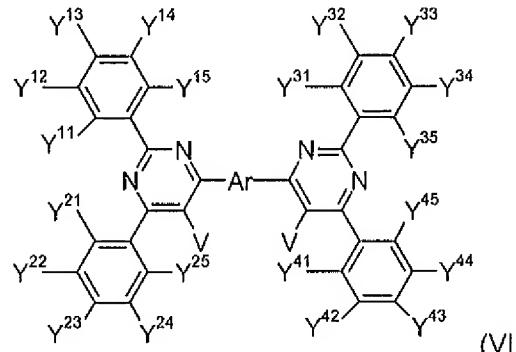
(IV), or

wherein V, W¹ to W⁵, X¹ to X⁵ and Y¹ to Y⁵ are as defined above, preferably W³, X³ and Y³

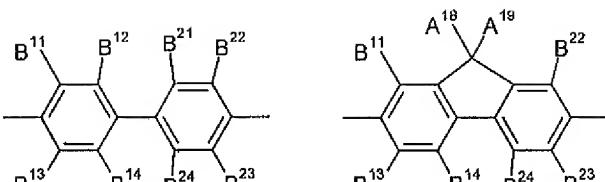
5 are selected from the group consisting of C₆-C₂₄aryl; C₆-C₂₄aryl which is substituted by G; C₂-C₂₄heteroaryl; C₂-C₂₄heteroaryl which is substituted by L, C₁-C₁₈alkoxy, -SR⁵; -NR⁵R⁶, wherein G, L, R⁵ and R⁶ are as defined above, V is H and/or W¹ and W⁵, Y¹ and Y⁵ as well as X¹ and X⁵ are independently of each other H; C₁-C₁₈alkyl; or C₁-C₁₈alkyl which is substituted by E and/or interrupted by D, wherein E and D are as defined above; or



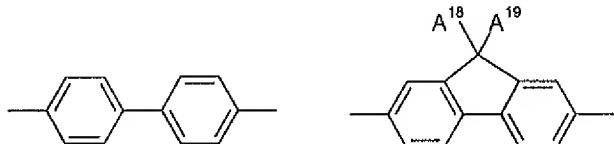
(VI), or



(VII), wherein



Ar is a group of formula B^{13} B^{19} B^{24} B^{23} , or B^{13} B^{19} B^{24} B^{23} , especially



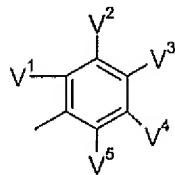
, or

W¹¹ to W¹⁵, W²¹ to W²⁵, W³¹ to W³⁵, W⁴¹ to W⁴⁵, Y¹¹ to Y¹⁵, Y²¹ to Y²⁵, Y³¹ to Y³⁵ and Y⁴¹ to Y⁴⁵ are independently of each other H; C₆-C₂₄aryl; C₆-C₂₄aryl which is substituted by G; C₁-

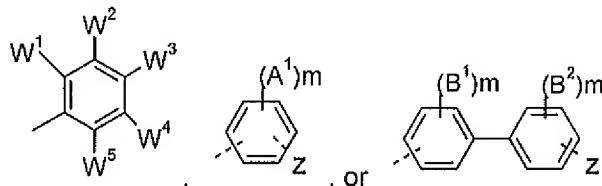
5 $C_{18}\text{alkyl}$; $C_1\text{-}C_{18}\text{alkyl}$ which is substituted by E and/or interrupted by D; $C_7\text{-}C_{18}\text{alkylaryl}$; $C_7\text{-}C_{18}\text{alkylaryl}$ which is substituted by E and/or interrupted by D; $C_2\text{-}C_{18}\text{alkenyl}$; $C_2\text{-}C_{18}\text{alkenyl}$ which is substituted by E and/or interrupted by D; $C_2\text{-}C_{18}\text{alkynyl}$; $C_2\text{-}C_{18}\text{alkynyl}$ which is substituted by E and/or interrupted by D; $C_1\text{-}C_{18}\text{alkoxy}$, $C_1\text{-}C_{18}\text{alkoxy}$ which is substituted by E and/or interrupted by D; $-\text{SR}^5$; $-\text{NR}^5\text{R}^6$; $C_2\text{-}C_{24}\text{heteroaryl}$; $C_2\text{-}C_{24}\text{heteroaryl}$ which is substituted by E and/or interrupted by D; $-\text{SOR}^4$; $-\text{SO}_2\text{R}^4$; $-\text{COR}^8$; $-\text{COOR}^7$; $-\text{CONR}^5\text{R}^6$; $C_4\text{-}C_{18}\text{cycloalkyl}$; $C_4\text{-}C_{18}\text{cycloalkyl}$ which is substituted by E and/or interrupted by D; $C_4\text{-}C_{18}\text{cycloalkenyl}$; $C_4\text{-}C_{18}\text{cycloalkenyl}$ which is substituted by E and/or interrupted by D; V is H; $C_6\text{-}C_{24}\text{aryl}$; $C_6\text{-}C_{24}\text{aryl}$ which is substituted by G; $C_1\text{-}C_{18}\text{alkyl}$; $C_1\text{-}C_{18}\text{alkyl}$ which is substituted by E and/or interrupted by D; $C_7\text{-}C_{18}\text{alkylaryl}$; $C_7\text{-}C_{18}\text{alkylaryl}$ which is substituted by E and/or interrupted by D; $C_2\text{-}C_{18}\text{alkenyl}$; $C_2\text{-}C_{18}\text{alkenyl}$ which is substituted by E and/or interrupted by D; $C_2\text{-}C_{18}\text{alkynyl}$; $C_2\text{-}C_{18}\text{alkynyl}$ which is substituted by E and/or interrupted by D; $C_1\text{-}C_{18}\text{alkoxy}$, $C_1\text{-}C_{18}\text{alkoxy}$ which is substituted by E and/or interrupted by D; $-\text{SR}^5$; or $-\text{NR}^5\text{R}^6$; $C_2\text{-}C_{24}\text{heteroaryl}$; $C_2\text{-}C_{24}\text{heteroaryl}$ which is substituted by L; $-\text{SOR}^4$; $-\text{SO}_2\text{R}^4$; $-\text{COR}^8$; $-\text{COOR}^7$; $-\text{CONR}^5\text{R}^6$; $C_4\text{-}C_{18}\text{cycloalkyl}$; $C_4\text{-}C_{18}\text{cycloalkyl}$ which is substituted by E and/or interrupted by D; $C_4\text{-}C_{18}\text{cycloalkenyl}$; $C_4\text{-}C_{18}\text{cycloalkenyl}$ which is substituted by E and/or interrupted by D; A^{18} and A^{19} are independently of each other H, $C_1\text{-}C_{18}\text{alkyl}$; $C_1\text{-}C_{18}\text{alkyl}$ which is substituted by E and/or interrupted by D; $C_6\text{-}C_{18}\text{aryl}$; $C_6\text{-}C_{18}\text{aryl}$ which is substituted by E, B^{11} to B^{14} and B^{21} to B^{24} are independently of each other H; $C_6\text{-}C_{18}\text{aryl}$; $C_6\text{-}C_{18}\text{aryl}$ which is substituted by G; $C_1\text{-}C_{18}\text{alkyl}$; $C_1\text{-}C_{18}\text{alkyl}$ which is substituted by E and/or interrupted by D; $C_7\text{-}C_{18}\text{alkylaryl}$; $C_7\text{-}C_{18}\text{alkylaryl}$ which is substituted by E and/or interrupted by D; $C_2\text{-}C_{18}\text{alkenyl}$; $C_2\text{-}C_{18}\text{alkenyl}$ which is substituted by E and/or interrupted by D; $C_2\text{-}C_{18}\text{alkynyl}$; $C_2\text{-}C_{18}\text{alkynyl}$ which is substituted by E and/or interrupted by D; $C_1\text{-}C_{18}\text{alkoxy}$, $C_1\text{-}C_{18}\text{alkoxy}$ which is substituted by E and/or interrupted by D; $-\text{SR}^5$; $-\text{NR}^5\text{R}^6$; $C_2\text{-}C_{18}\text{heteroaryl}$; $C_2\text{-}C_{18}\text{heteroaryl}$

C₁₈heteroaryl which is substituted by L; -SOR⁴; -SO₂R⁴; -COR⁸; -COOR⁷; or -CONR⁵R⁶; C₄-C₁₈cycloalkyl; C₄-C₁₈cycloalkyl which is substituted by E and/or interrupted by D; C₄-C₁₈cycloalkenyl; C₄-C₁₈cycloalkenyl which is substituted by E and/or interrupted by D; wherein D, E, G, L, R⁴, R⁵, R⁶, R⁷ and R⁸ are as defined above, and V in formula VI or VII is 5 preferably H.

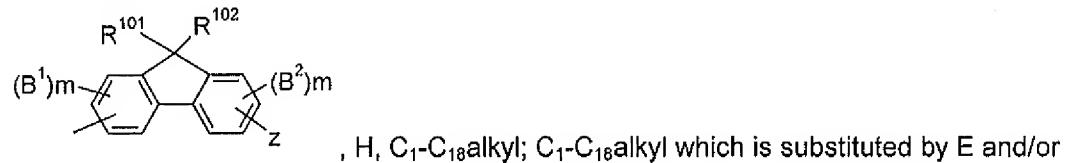
Moreover, pyrimidine compounds of formula I are preferred, wherein V is a group of the



formula , H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl, C₂-C₁₈alkenyl which is substituted by E and/or interrupted by 10 D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy; C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; or -NR⁵R⁶; and/or



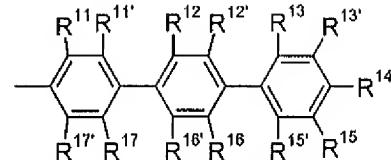
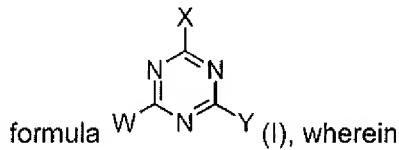
W is a group of the formula



15 , H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl, C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy; C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; or -NR⁵R⁶; wherein W¹ to W⁵, D, V¹ to V⁵, E, A¹, B¹, B², R⁵, R⁶, m and Z are as defined above and R¹⁰¹ and R¹⁰² are independently of each other H, C₁-C₈alkyl, C₆-C₂₄aryl, or C₅-C₇cycloalkyl, in particular H or C₁₋₄-alkyl.

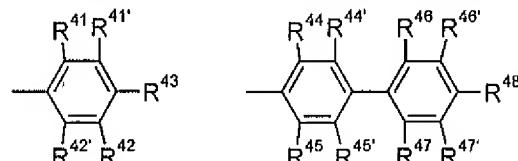
The compounds of general formula IV, VI and VII are novel and form a further subject of the present application.

In one preferred embodiment the present invention is directed to pyrimidine compounds of

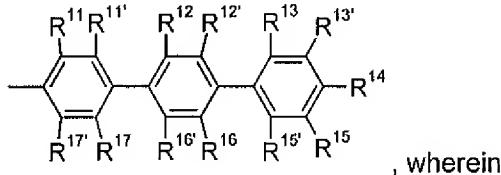


at least one of the groups W, X and Y is a group of formula

and the other groups are independently of each other an aryl group or a heteroaryl group,



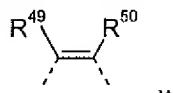
5 especially a group of formula



$\text{R}^{11}, \text{R}^{11'}, \text{R}^{12}, \text{R}^{12'}, \text{R}^{13}, \text{R}^{13'}, \text{R}^{15}, \text{R}^{15'}, \text{R}^{16}, \text{R}^{16'}, \text{R}^{17}, \text{R}^{17'}, \text{R}^{41}, \text{R}^{41'}, \text{R}^{42}, \text{R}^{42'}, \text{R}^{44}, \text{R}^{44'}, \text{R}^{45}, \text{R}^{45'}, \text{R}^{46}, \text{R}^{46'}, \text{R}^{47}$ and $\text{R}^{47'}$ are independently of each other H, E, $\text{C}_6\text{-C}_{18}$ aryl; $\text{C}_6\text{-C}_{18}$ aryl which is substituted by E; $\text{C}_1\text{-C}_{18}$ alkyl; $\text{C}_1\text{-C}_{18}$ alkyl which is substituted by E and/or interrupted by D;

10 $\text{C}_7\text{-C}_{18}$ aralkyl; or $\text{C}_7\text{-C}_{18}$ aralkyl which is substituted by E; or

$\text{R}^{11'}$ and $\text{R}^{12}, \text{R}^{12'}$ and $\text{R}^{13}, \text{R}^{13'}$ and $\text{R}^{16}, \text{R}^{16'}$ and $\text{R}^{17}, \text{R}^{17'}$, $\text{R}^{44'}$ and R^{46} and/or $\text{R}^{45'}$ and R^{47} are each a divalent group L^1 selected from an oxygen atom, an sulfur atom, $>\text{CR}^{118}\text{R}^{119}>\text{SiR}^{118}\text{R}^{119}$, or

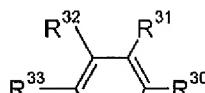


, wherein

R^{118} and R^{119} are independently of each other $\text{C}_1\text{-C}_{18}$ alkyl; $\text{C}_1\text{-C}_{18}$ alkoxy, $\text{C}_6\text{-C}_{18}$ aryl; $\text{C}_7\text{-C}_{18}$ aralkyl;

15

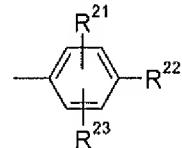
R^{11} and $\text{R}^{11'}, \text{R}^{12}$ and $\text{R}^{12'}, \text{R}^{13}$ and $\text{R}^{13'}, \text{R}^{13'}$ and $\text{R}^{14}, \text{R}^{14}$ and $\text{R}^{15}, \text{R}^{15}$ and $\text{R}^{15'}, \text{R}^{16}$ and $\text{R}^{16'}, \text{R}^{17}$ and $\text{R}^{17'}, \text{R}^{41}$ and $\text{R}^{41'}, \text{R}^{42}$ and $\text{R}^{42'}, \text{R}^{42'}$ and $\text{R}^{43}, \text{R}^{41'}$ and $\text{R}^{43}, \text{R}^{44}$ and $\text{R}^{44'}, \text{R}^{45}$ and $\text{R}^{45'}, \text{R}^{46}$ and $\text{R}^{46'}, \text{R}^{47}$ and $\text{R}^{47'}, \text{R}^{46'}$ and R^{48} and/or $\text{R}^{47'}$ and R^{48} are each a divalent group



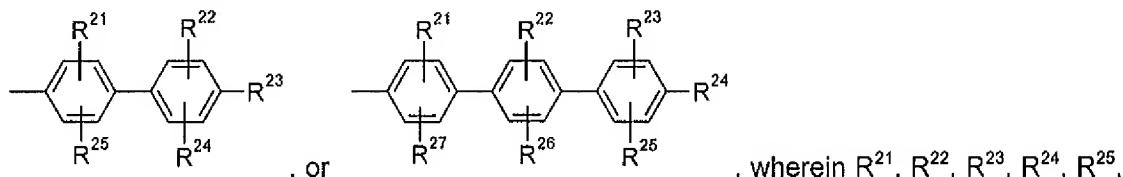
, wherein

R^{30} , R^{31} , R^{32} , R^{33} , R^{49} and R^{50} are independently of each other H, C_1 - C_{18} alkyl; C_1 - C_{18} alkyl, which is substituted by E and/or interrupted by D; E; C_6 - C_{18} aryl; C_6 - C_{18} aryl, which is substituted by E;

R^{14} is H, C_2 - C_{30} heteroaryl, C_6 - C_{30} aryl, or C_6 - C_{30} aryl which is substituted by E, C_1 - C_{18} alkyl; or



5 C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D; especially

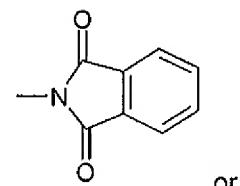


R^{26} and R^{27} are independently of each other H, E, C_1 - C_{18} alkyl; C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D; E; C_7 - C_{18} aralkyl; C_7 - C_{18} aralkyl which is substituted by E;

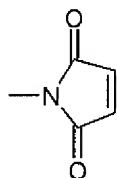
10 R^{43} and R^{48} are independently of each other H, E; C_1 - C_{18} alkyl; C_1 - C_{18} alkyl, which is substituted by E and/or interrupted by D; C_2 - C_{30} heteroaryl; C_7 - C_{18} aralkyl; C_7 - C_{18} aralkyl which is substituted by E;

D is -CO-; -COO-; -OCOO-; -S-; -SO-; -SO₂-; -O-; -NR⁵-; SiR⁵R⁶-; -POR⁵-; -CR⁹=CR¹⁰-; or -C≡C-;

15 E is -OR⁵; -SR⁵; -NR⁵R⁶; -COR⁸; -COOR⁷; -CONR⁵R⁶; -CN; or halogen, especially F, or Cl; wherein R⁵ and R⁶ are independently of each other C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by C_1 - C_{18} alkyl, C_1 - C_{18} alkyl; or C_1 - C_{18} alkyl which is interrupted by -O-; or



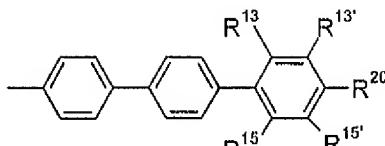
R^5 and R^6 together form a five or six membered ring, in particular



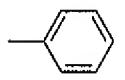
20 R^7 is C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by C_1 - C_{18} alkyl, C_1 - C_{18} alkyl; or C_1 - C_{18} alkyl which is interrupted by -O-;

R^8 is C_7 - C_{12} alkylaryl; C_1 - C_{18} alkyl; or C_1 - C_{18} alkyl which is interrupted by -O-; and

R^9 and R^{10} are independently of each other H, C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by C_1 - C_{18} alkyl, C_1 - C_{18} alkyl; or C_1 - C_{18} alkyl which is interrupted by -O-.



W is preferably a group of formula

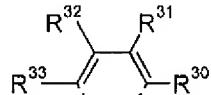


R¹³, R^{13'}, R¹⁵ and R^{15'} are H and R²⁰ is H, especially

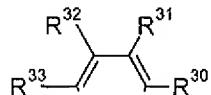
, or

R¹³ and R¹⁵ are H, R^{13'} and R^{15'} are independently of each other H, C₁-C₈alkyl, or C₁-C₈alkoxy,

5 and R²⁰ is H, C₁-C₈alkyl, or C₁-C₈alkoxy; or



R¹³, R¹⁵ and R^{15'} are H, and R^{13'} and R²⁰ are



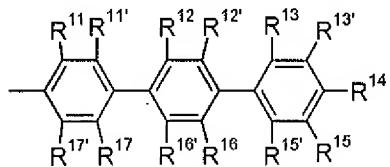
R²⁰, R¹⁵ and R^{15'} are H, and R¹³ and R^{13'} are

wherein

R³⁰, R³¹, R³² and R³³ are H, C₁-C₈alkyl, or C₁-C₈alkoxy, and

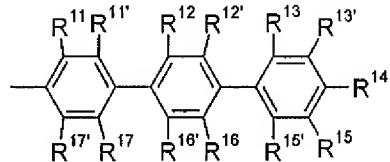
10 X and Y are as defined above.

According to the present invention at least one of the groups W, X and Y, preferably two of the groups W, X and Y, most preferred all three groups W, X and Y are a group of formula

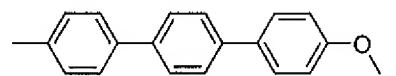
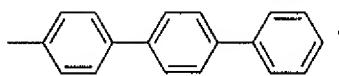


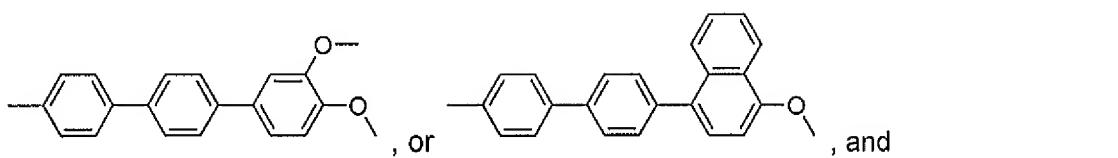
15

Accordingly, in one preferred embodiment of the present invention the pyrimidine compound is a compound of formula I, wherein W and Y or W and X (= X and Y) are independently of

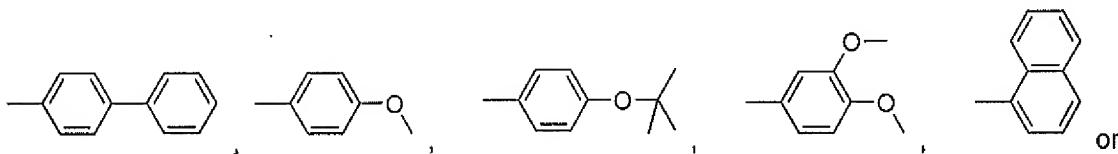


each other a group of formula





and
 $\text{R}^{41}\text{R}^{41'}$
 $\text{R}^{42}\text{R}^{42'}$
 R^{43}
 $\text{R}^{44}\text{R}^{44'}$
 $\text{R}^{45}\text{R}^{45'}$
 $\text{R}^{46}\text{R}^{46'}$
 $\text{R}^{47}\text{R}^{47'}$
 R^{48}
X is a group of formula , or , especially ,



5 , wherein R^{11} , $\text{R}^{11'}$, R^{12} , $\text{R}^{12'}$, R^{13} , $\text{R}^{13'}$, R^{15} , $\text{R}^{15'}$, R^{16} , $\text{R}^{16'}$, R^{17} and $\text{R}^{17'}$, R^{41} , $\text{R}^{41'}$, R^{42} , $\text{R}^{42'}$, R^{44} , $\text{R}^{44'}$, R^{45} , $\text{R}^{45'}$, R^{46} , $\text{R}^{46'}$, R^{47} , $\text{R}^{47'}$, R^{48} and $\text{R}^{48'}$ are as defined above, especially H, C₁-C₈alkyl, C₁-C₈alkoxy, or phenyl.

10 R^{11} , $\text{R}^{11'}$, R^{12} , $\text{R}^{12'}$, R^{13} , $\text{R}^{13'}$, R^{15} , $\text{R}^{15'}$, R^{16} , $\text{R}^{16'}$, R^{17} and $\text{R}^{17'}$, R^{41} , $\text{R}^{41'}$, R^{42} , $\text{R}^{42'}$, R^{44} , $\text{R}^{44'}$, R^{45} ,
 $\text{R}^{45'}$, R^{46} , $\text{R}^{46'}$, R^{47} , and $\text{R}^{47'}$ as well as R^{14} , R^{43} , and R^{48} are preferably independently of each other H, E; or C₁-C₈alkyl, especially H, C₁-C₄alkyl, C₁-C₄alkoxy, or phenyl; wherein E is -OR⁵; -SR⁵; -NR⁵R⁶; -COR⁸; -COOR⁷; -CONR⁵R⁶; -CN; -OCOOR⁷; or halogen, especially F; wherein R⁵ and R⁶ are independently of each other C₆-C₁₂aryl, or C₁-C₈alkyl;

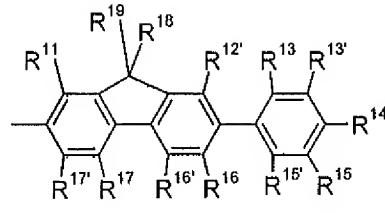
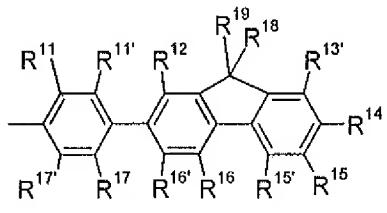
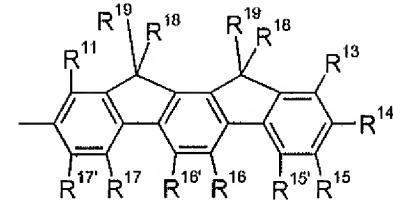
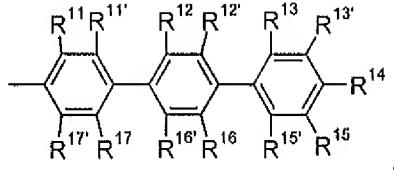
R^7 is C₇-C₁₂ alkylaryl, or C₁-C₈alkyl; and

15 R^8 is C₆-C₁₂aryl; or C₁-C₈alkyl, or
 R^{11} and $\text{R}^{11'}$, R^{12} and $\text{R}^{12'}$, R^{13} and $\text{R}^{13'}$, $\text{R}^{13'}$ and R^{14} , R^{41} and $\text{R}^{41'}$ and R^{43} , R^{44} and $\text{R}^{44'}$,

R^{46} and $\text{R}^{46'}$, $\text{R}^{46'}$ and R^{48} and/or R^{47} and R^{48} are each a divalent group



20 In one more preferred embodiment of the present invention W, X and Y are independently of each other a group of formula



, wherein

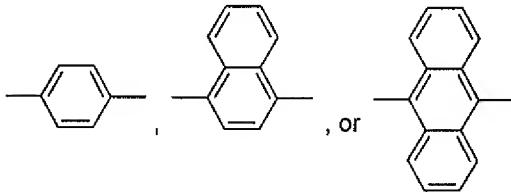
R¹¹, R^{11'}, R¹², R^{12'}, R¹³, R^{13'}, R¹⁵, R^{15'}, R¹⁶, R^{16'}, R¹⁷ and R^{17'} are independently of each other H, C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by E; E, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₇-C₁₈aralkyl; C₇-C₁₈aralkyl which is substituted by E; and R¹⁴, R¹⁸ and R¹⁹ are as defined above, or

W is a group of the formula -W¹-W²-W³,

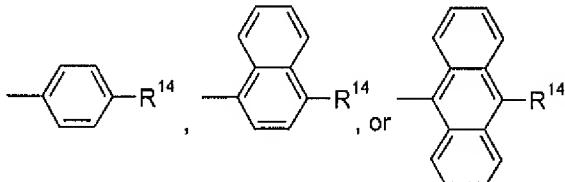
X is a group of the formula -X¹-X²-X³ and/or

Y is a group of the formula -Y¹-Y²-Y³, wherein W¹, W², X¹, X², Y¹ and Y² are independently of

each other a group of formula



and W³, X³ and Y³ are independently of each



other a group of formula

defined above.

wherein R¹⁴ is as

15 W, X and Y can be different, but have preferably the same meaning.

Pyrimidine compounds of formula I are preferred, wherein R¹¹, R^{11'}, R¹², R^{12'}, R¹³, R^{13'}, R¹⁵, R^{15'}, R¹⁶, R^{16'}, R¹⁷ and R^{17'}, R⁴¹, R^{41'}, R⁴², R^{42'}, R⁴⁴, R^{44'}, R⁴⁵, R^{45'}, R⁴⁶, R^{46'}, R⁴⁷, and R^{47'} are independently of each other H, E; or C₁-C₈alkyl; wherein

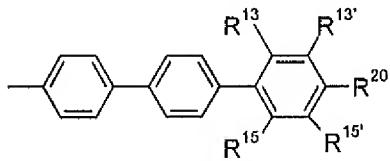
20 E is -OR⁵; -SR⁵; -NR⁵R⁶; -COR⁸; -COOR⁷; -CONR⁵R⁶; -CN; -OCOOR⁷; or halogen; wherein R⁵ and R⁸ are independently of each other C₆-C₁₂aryl; or C₁-C₈alkyl;

R^7 is C_7 - C_{12} alkylaryl, or C_1 - C_8 alkyl; and

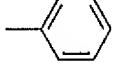
R^8 is C_6 - C_{12} aryl, or C_1 - C_8 alkyl.

Especially preferred are pyrimidine compounds of formula I, wherein

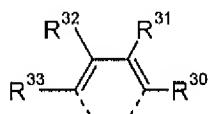
5 W, X and Y are independently of each other a group of formula



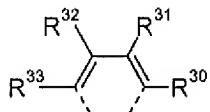
, wherein R^{13} , $R^{13'}$, R^{15} and $R^{15'}$ are H and R^{20} is H,

especially  , or

R^{13} and R^{15} are H, $R^{13'}$ and $R^{15'}$ are independently of each other H, C_1 - C_8 alkyl, or C_1 - C_8 alkoxy, and R^{20} is H, C_1 - C_8 alkyl, or C_1 - C_8 alkoxy; or



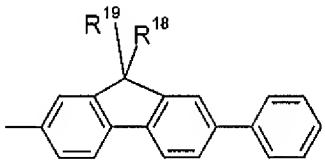
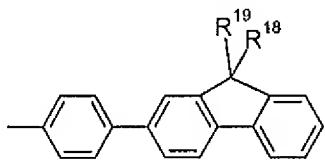
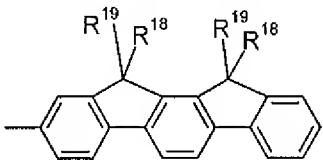
10 R^{13} , R^{15} and $R^{15'}$ are H, and $R^{13'}$ and R^{20} are , or



R^{20} , R^{15} and $R^{15'}$ are H, and R^{13} and $R^{13'}$ are , wherein

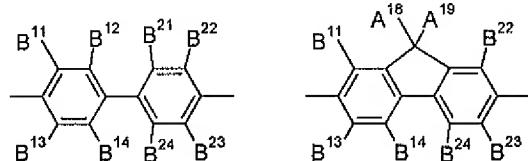
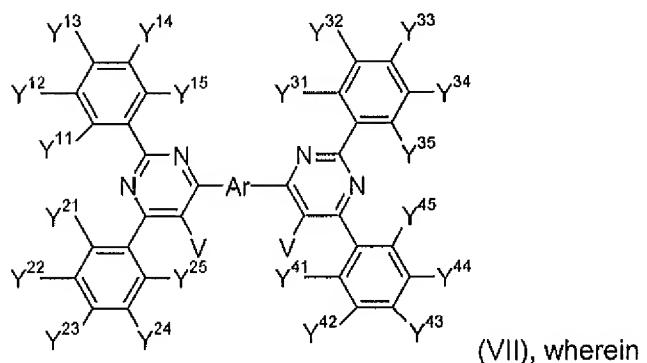
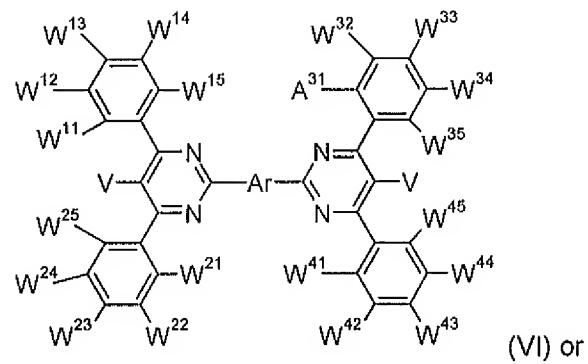
R^{30} , R^{31} , R^{32} and R^{33} are H, C_1 - C_8 alkyl, or C_1 - C_8 alkoxy; or wherein

W, X and Y are independently of each other a group of formula

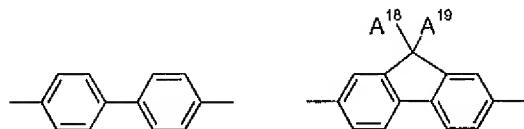


15 wherein R^{18} and R^{19} are independently of each other C_1 - C_8 alkyl.

In a further preferred embodiment the present invention is directed to pyrimidine compounds of formula



Ar is a group of formula , or , especially



, or ,

5 W^{11} to W^{15} , W^{21} to W^{25} , W^{31} to W^{35} , W^{41} to W^{45} , Y^{11} to Y^{15} , Y^{21} to Y^{25} , Y^{31} to Y^{35} and Y^{41} to Y^{45} are independently of each other H; C_6 - C_{24} aryl; C_6 - C_{24} aryl which is substituted by G; C_1 - C_{18} alkyl; C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D; C_7 - C_{18} alkylaryl; C_7 - C_{18} alkylaryl which is substituted by E and/or interrupted by D; C_2 - C_{18} alkenyl; C_2 - C_{18} alkenyl which is substituted by E and/or interrupted by D; C_2 - C_{18} alkynyl; C_2 - C_{18} alkynyl which is substituted by E and/or interrupted by D; C_1 - C_{18} alkoxy, C_1 - C_{18} alkoxy which is substituted by E and/or interrupted by D; $-SR^5$; $-NR^5R^6$; C_2 - C_{24} heteroaryl; C_2 - C_{24} heteroaryl which is substituted by L; $-SOR^4$; $-SO_2R^4$; $-COR^8$; $-COOR^7$; $-CONR^5R^6$; C_4 - C_{18} cycloalkyl; C_4 - C_{18} cycloalkyl which is substituted by E and/or interrupted by D; C_4 - C_{18} cycloalkenyl; C_4 - C_{18} cycloalkenyl which is substituted by E and/or interrupted by D;

10 C_1 - C_{18} alkoxy, C_1 - C_{18} alkoxy which is substituted by E and/or interrupted by D; $-SR^5$; $-NR^5R^6$; C_2 - C_{24} heteroaryl; C_2 - C_{24} heteroaryl which is substituted by L; $-SOR^4$; $-SO_2R^4$; $-COR^8$; $-COOR^7$; $-CONR^5R^6$; C_4 - C_{18} cycloalkyl; C_4 - C_{18} cycloalkyl which is substituted by E and/or interrupted by D; C_4 - C_{18} cycloalkenyl; C_4 - C_{18} cycloalkenyl which is substituted by E and/or interrupted by D;

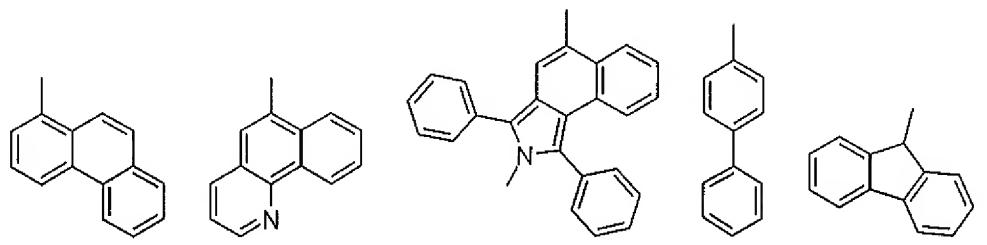
V is H; C₆-C₂₄aryl; C₆-C₂₄aryl which is substituted by G; C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₇-C₁₈alkylaryl; C₇-C₁₈alkylaryl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl; C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D.

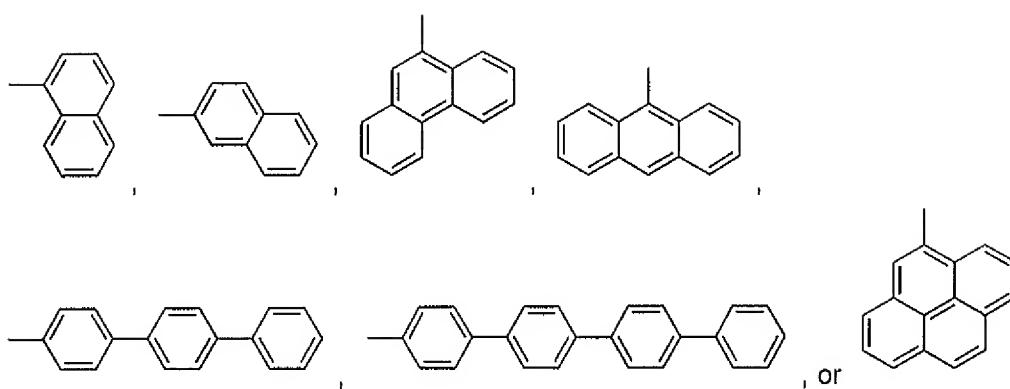
5 D; C₁-C₁₈alkoxy, C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; or -NR⁵R⁶; C₂-C₂₄heteroaryl; C₂-C₂₄heteroaryl which is substituted by L; -SOR⁴; -SO₂R⁴; -COR⁸; -COOR⁷; -CONR⁵R⁶; C₄-C₁₈cycloalkyl; C₄-C₁₈cycloalkyl which is substituted by E and/or interrupted by D; C₄-C₁₈cycloalkenyl; C₄-C₁₈cycloalkenyl which is substituted by E and/or interrupted by D;

10 A¹⁸ and A¹⁹ are independently of each other H, C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by E, B¹¹ to B¹⁴, B²¹ to B²⁴, B³¹ to B³⁴ and B⁴¹ to B⁴⁴ are independently of each other H; C₆-C₁₈aryl; C₆-C₁₈aryl which is substituted by G; C₁-C₁₈alkyl; C₁-C₁₈alkyl which is substituted by E and/or interrupted by D; C₇-C₁₈alkylaryl; C₇-C₁₈alkylaryl which is substituted by E and/or interrupted by D; C₂-C₁₈alkenyl; C₂-C₁₈alkenyl which is substituted by E and/or interrupted by D; C₂-C₁₈alkynyl; C₂-C₁₈alkynyl which is substituted by E and/or interrupted by D; C₁-C₁₈alkoxy, C₁-C₁₈alkoxy which is substituted by E and/or interrupted by D; -SR⁵; -NR⁵R⁶; C₂-C₁₈heteroaryl; C₂-C₁₈heteroaryl which is substituted by L; -SOR⁴; -SO₂R⁴; -COR⁸; -COOR⁷; or -CONR⁵R⁶; C₄-C₁₈cycloalkyl; C₄-C₁₈cycloalkyl which is substituted by E and/or interrupted by D; C₄-C₁₈cycloalkenyl; C₄-C₁₈cycloalkenyl which is substituted by E and/or interrupted by D;

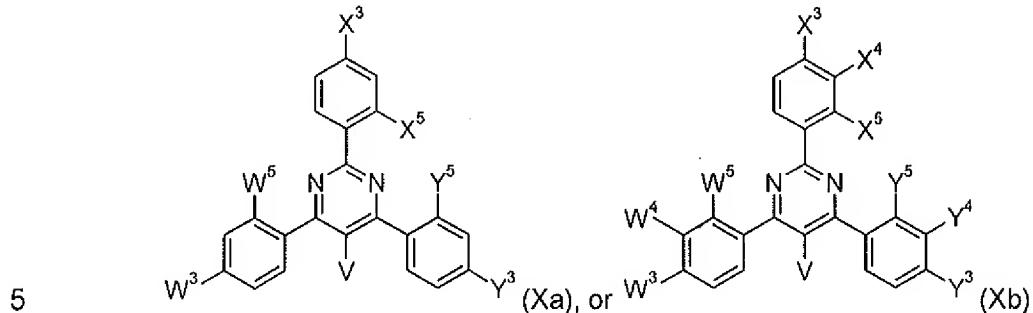
15 wherein D, E, G, L, R⁴, R⁵, R⁶, R⁷ and R⁸ are as defined above.

In a preferred embodiment W and Y are groups of the formula





Particularly preferred are pyrimidine compounds of the following formula:



wherein V is H, or C₁-C₆-alkyl,

X³ and X⁴ are independently of each other H, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆thioalkyl, or phenyl,

X⁵ is H, or C₁-C₆alkoxy,

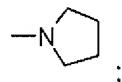
10 W⁵ is H, C₁-C₆alkyl, or O(CH₂)_{n1}-X,

Y⁵ is H, C₁-C₆alkyl, or O(CH₂)_{n1}-X,

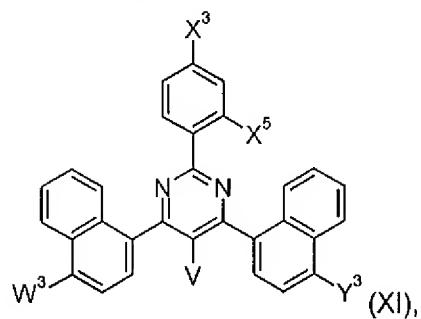
Y³, Y⁴, W³ and W⁴ are independently of each other C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆thioalkyl, halogen, in particular Br, phenyl, or O(CH₂)_{n1}-X, wherein n1 is an integer of 1 to 5 and X is -

O-(CH₂)_{m1}CH₃, -OC(O)-(CH₂)_{m1}CH₃, -C(O)-O-C₁-C₆alkyl, -NR¹⁰³R¹⁰⁴, wherein m1 is an

15 integer of 0 to 5 and R¹⁰³ and R¹⁰⁴ are independently of each other H, or C₁-C₆-alkyl, or R¹⁰³



and R¹⁰⁴ together form a five or six membered heterocyclic ring, in particular



wherein V is H, or C₁-C₈alkyl,

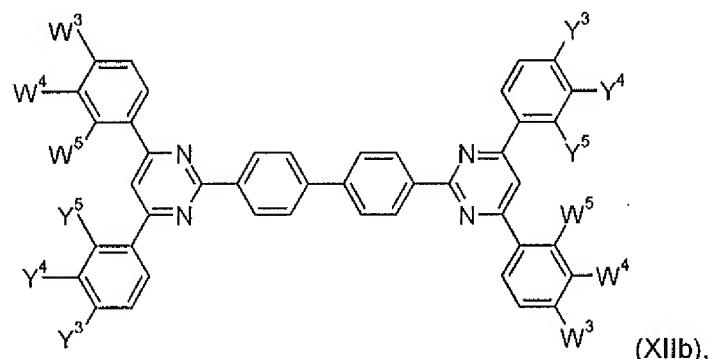
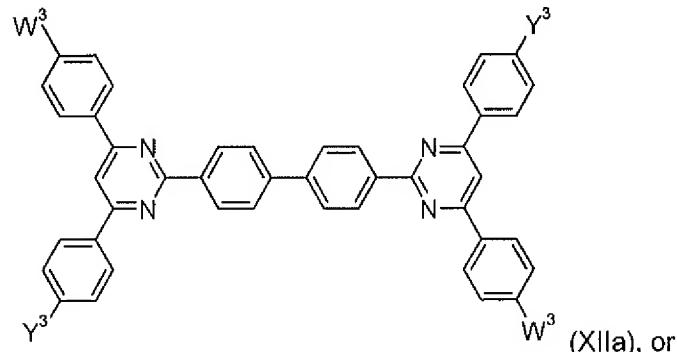
W³ is H, C₁-C₈alkyl, or C₁-C₈alkoxy,

X³ is H, C₁-C₈alkoxy, phenyl or O(CH₂)_{n1}-X,

X⁵ is H, C₁-C₈alkoxy, phenyl or O(CH₂)_{n1}-X,

5 Y³ is H, C₁-C₈alkyl, or C₁-C₈alkoxy, wherein n1 is an integer of 1 to 4 and X is -

O-(CH₂)_{m1}CH₃, -OC(O)-(CH₂)_{m1}CH₃, -C(O)-O-C₁-C₈alkyl, wherein m1 is an integer of 0 to 5;



wherein W³ and W⁴ are independently of each other H, -NR¹⁰³R¹⁰⁴, C₁-C₈thioalkyl, or C₁-

10 C₈alkoxy,

Y³ and Y⁴ are independently of each other H, -NR¹⁰³R¹⁰⁴, C₁-C₈thioalkyl, or C₁-C₈alkoxy,

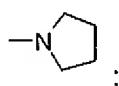
wherein R¹⁰³ and R¹⁰⁴ are independently of each other H, or C₁-C₈alkyl.

W⁵ is H, C₁-C₈alkyl, or O(CH₂)_{n1}-X,

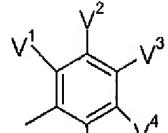
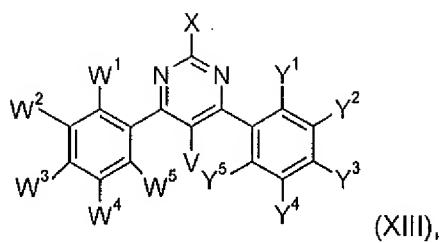
Y⁵ is H, C₁-C₈alkyl, or O(CH₂)_{n1}-X,

15 wherein n1 is an integer of 1 to 5 and X is -O-(CH₂)_{m1}CH₃, -OC(O)-(CH₂)_{m1}CH₃, -C(O)-O-C₁-

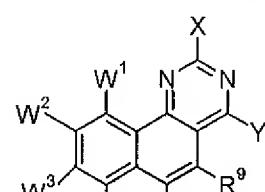
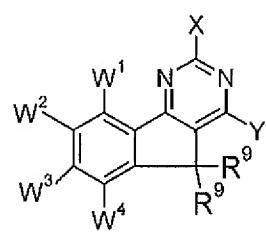
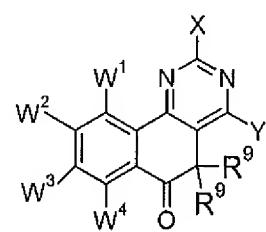
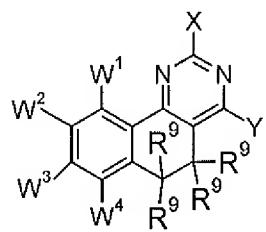
C₈alkyl, -NR¹⁰³R¹⁰⁴, wherein m1 is an integer of 0 to 5 and R¹⁰³ and R¹⁰⁴ are independently of each other H, or C₁-C₈-alkyl, or R¹⁰³ and R¹⁰⁴ together form a five or six membered



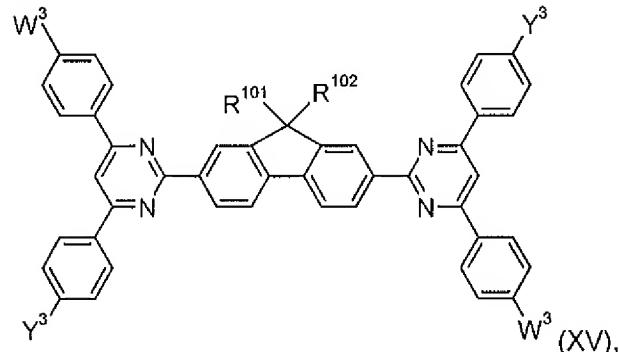
heterocyclic ring, in particular



wherein V, W¹ to W⁵, X, Y¹ to Y⁵ are as defined above, wherein V is preferably with the proviso that at least one substituent V¹ to V⁵ is different from hydrogen;



(XIVe), wherein X, Y, W¹ to W⁴, and R⁹ are as defined above;

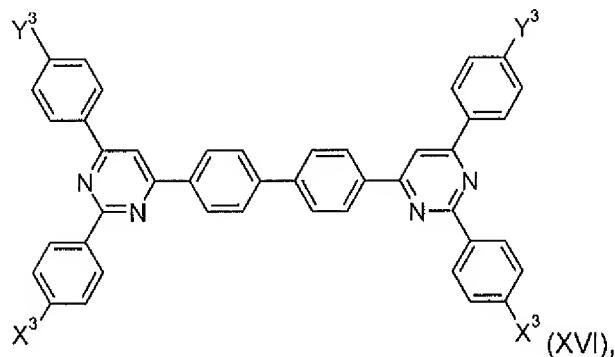


wherein W³ is H, -NR¹⁰³R¹⁰⁴, C₁-C₈thioalkyl, or C₁-C₈alkoxy,

Y³ is H, -NR¹⁰³R¹⁰⁴, C₁-C₈thioalkyl, or C₁-C₈alkoxy, wherein R¹⁰³ and R¹⁰⁴ are independently

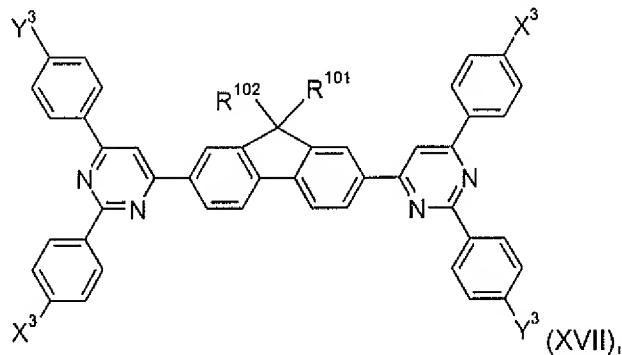
of each other H, or C₁-C₈alkyl,

R¹⁰¹ and R¹⁰² are independently of each other H, C₁-C₈alkyl, phenyl, or C₆-C₇cycloalkyl, in particular cyclohexyl;



wherein Y^3 is H, $-NR^{103}R^{104}$, C_1 - C_8 thioalkyl, or C_1 - C_8 alkoxy,

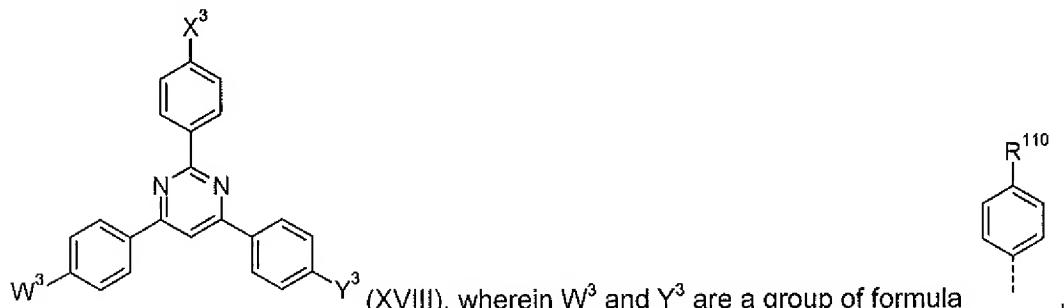
X^3 is H, $-NR^{103}R^{104}$, C_1 - C_8 thioalkyl, or C_1 - C_8 alkoxy, wherein R^{103} and R^{104} are independently of each other H, or C_1 - C_8 alkyl;



Y^3 is H, $-NR^{103}R^{104}$, C_1 - C_8 thioalkyl, or C_1 - C_8 alkoxy,

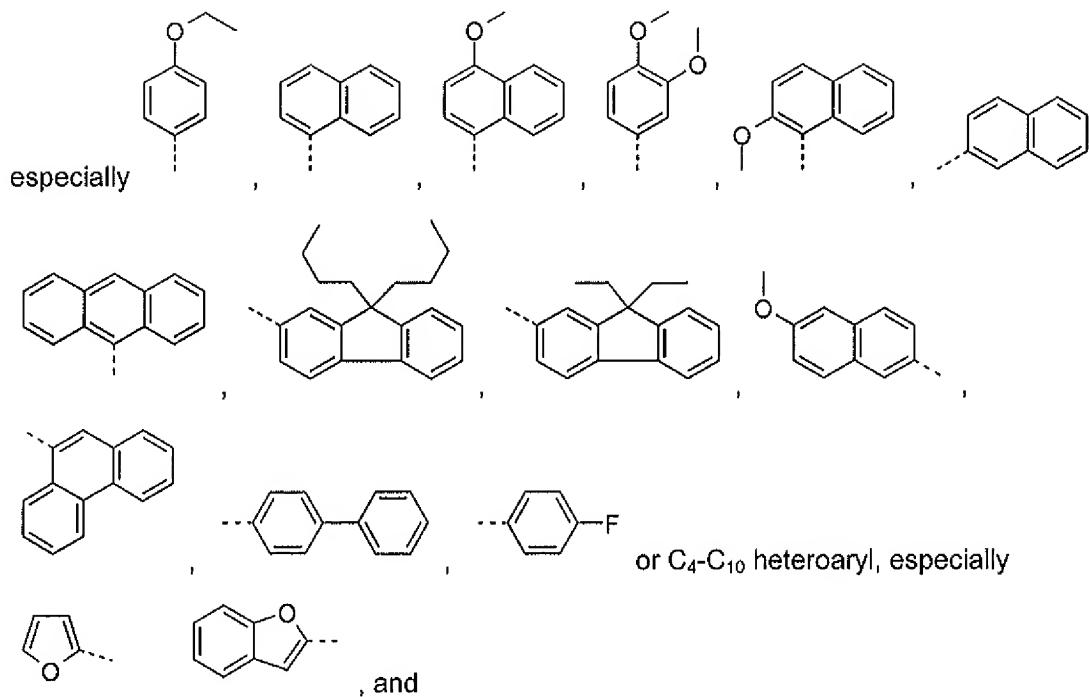
X^3 is H, $-NR^{103}R^{104}$, C_1 - C_8 thioalkyl, or C_1 - C_8 alkoxy, wherein R^{103} and R^{104} are independently of each other H, or C_1 - C_8 alkyl, R^{101} and R^{102} are independently of each other H, C_1 - C_8 alkyl, phenyl, or C_6 - C_7 cycloalkyl, in particular cyclohexyl;

10 pyrimidine compounds of formula

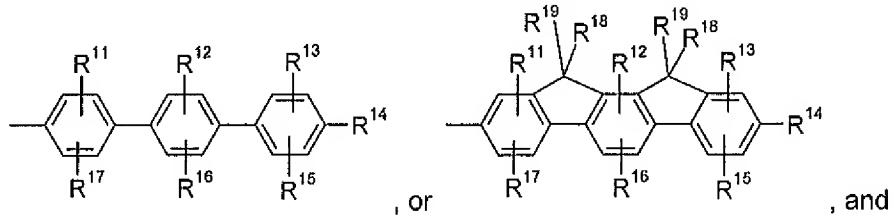


wherein

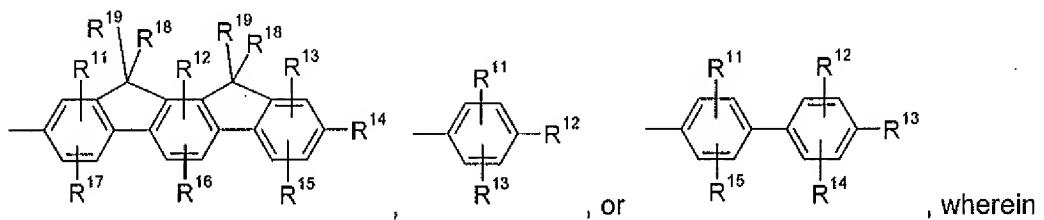
R^{110} is C_6 - C_{10} -aryl, such as phenyl, 1-naphthyl, 2-naphthyl, 3- or 4-biphenyl, 9-phenanthryl, 2- or 9-fluorenyl, which is optionally substituted by C_1 - C_6 -alkyl, or C_1 - C_4 -alkoxy,



5 X^3 is H, C₁-C₆-alkyl, C₁-C₄-alkoxy, Ph, or ; and
pyrimidine compounds of formula I, wherein
V is hydrogen,
W and Y are independently of each other a group of formula



10 X is a group of formula R¹¹ R¹⁸ R¹⁵



R^{11} , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} and R^{17} are independently of each other H, C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by E; E, C_1 - C_{18} alkyl; C_1 - C_{18} alkyl which is substituted by E and/or interrupted by D; C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by E;

R^{18} and R^{19} are independently of each other H, C_1 - C_{18} alkyl; C_1 - C_{18} alkyl which is substituted

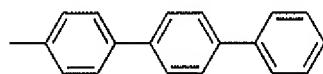
5 by E and/or interrupted by D; C_6 - C_{18} aryl; C_6 - C_{18} aryl which is substituted by E;

D is -CO-; -COO-; -OCOO-; -S-; -SO-; -SO₂-; -O-; -NR⁵-; -SiR⁶R⁶-; -POR⁶-; -CR⁶=CR⁶-; or -C≡C-;

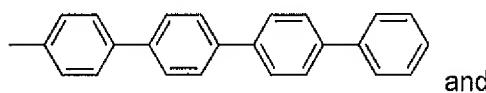
E is -OR⁵; -SR⁵; -NR⁵R⁶; -COR⁸; -COOR⁷; -CONR⁵R⁶; -CN; -OCOOR⁷; or halogen; wherein

10 R^5 , R^6 , R^7 and R^8 are as defined above, wherein pyrimidine compounds of formula I, wherein

V is hydrogen,



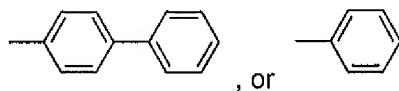
W and Y are a group of formula



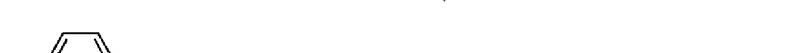
and



X is a group of formula

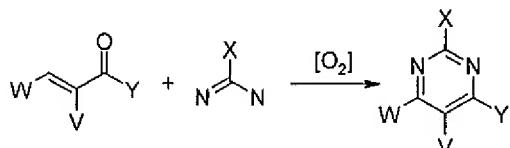


, or



15 The present pyrimidine compounds are known or can be prepared according to or analogous to known procedures. The present pyrimidine compounds are for instance derivatives of known hydroxyphenyl pyrimidine compounds: US-A-3,442,898, US-A-5,597,854 and US-A-5,753,729, the relevant parts of which are hereby incorporated by reference. The present pyrimidine compounds can for instance be prepared according to or analogous to the

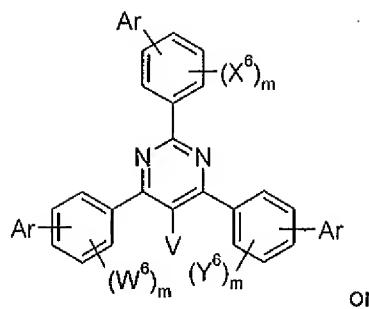
20 following procedures (Suzuki aryl-aryl cross coupling reaction: Chem. Commun., 2002, 874 – 875; DE-A-3001188, J. Org. Chem. Vol. 36, 1971, 3382-3385):



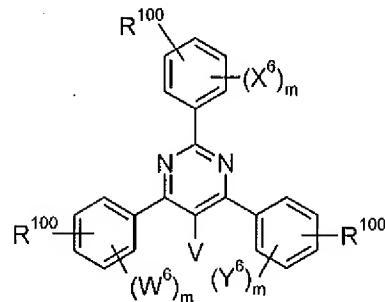
An amidine hydrochloride is added to a 2-propen-1-one derivative in a C_1 - C_4 -alcohol, like methanol, ethanol, propanol or butanol. A base, for example, an alkali metal hydroxide or

25 alkoxide, such as sodium methoxide or potassium hydroxide is added and the solution is stirred in the presence of oxygen, for example dry air.

The pyrimidine compounds of the present invention, comprising the following units:

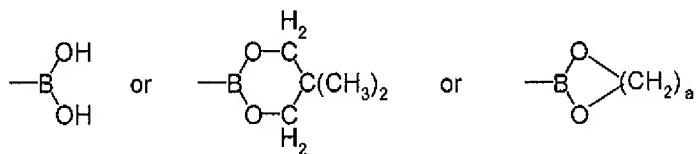


can be prepared according to a process, which comprises reacting a derivative of formula



wherein X^6 , V , W^6 , Y^6 and m are as defined above,

5 R^{100} stands for halogen such as chloro or bromo, preferably bromo, or E having the meaning of



wherein a is 2 or 3,

with boronic acid derivative

10 $E-Ar$,

or - in case R^{100} is not halogen -

$Hal-Ar$,

wherein Hal stands for halogen, preferably for bromo,

wherein Ar is C_6-C_{30} -aryl or C_2-C_{30} -heteroaryl, which can be substituted, in the presence of

15 an allylpalladium catalyst of the μ -halo(triisopropylphosphine)(η^3 -allyl)palladium(II) type (see for example WO99/47474).

Preferably, the reaction is carried out in the presence of an organic solvent, such as an aromatic hydrocarbon or a usual polar organic solvent, such as benzene, toluene, xylene, tetrahydrofuran, or dioxane, or mixtures thereof, most preferred toluene. Usually, the

20 amount of the solvent is chosen in the range of from 1 to 10 l per mol of boronic acid derivative. Also preferred, the reaction is carried out under an inert atmosphere such as nitrogen, or argon.

Further, it is preferred to carry out the reaction in the presence of an aqueous base, such as an alkali metal hydroxide or carbonate such as NaOH, KOH, Na₂CO₃, K₂CO₃, Cs₂CO₃ and the like, preferably an aqueous K₂CO₃ solution is chosen. Usually, the molar ratio of the base to compound III is chosen in the range of from 0.5:1 to 50:1.

5 Generally, the reaction temperature is chosen in the range of from 40 to 180°C, preferably under reflux conditions.

Preferred, the reaction time is chosen in the range of from 1 to 80 hours, more preferably from 20 to 72 hours.

In a preferred embodiment a usual catalyst for coupling reactions or for polycondensation

10 reactions is used, preferably Pd-based catalyst such as known tetrakis(triarylphosphonium)-palladium, preferably (Ph₃P)₄Pd and derivatives thereof. Usually, the catalyst is added in a molar ratio from inventive DPP polymer to the catalyst in the range of from 100:1 to 10:1, preferably from 50:1 to 30:1.

Also preferred, the catalyst is added as in solution or suspension. Preferably, an appropriate 15 organic solvent such as the ones described above, preferably benzene, toluene, xylene, THF, dioxane, more preferably toluene, or mixtures thereof, is used. The amount of solvent usually is chosen in the range of from 1 to 10 l per mol of boronic acid derivative.

The obtained inventive polymer can be isolated by well-known methods. Preferably, after cooling down the reaction mixture to room temperature, it is poured into acetone and the 20 obtained precipitation is filtered off, washed and dried.

C₁-C₁₈alkyl is a branched or unbranched radical such as for example methyl, ethyl, propyl, isopropyl, n-butyl, sec-butyl, isobutyl, tert-butyl, 2-ethylbutyl, n-pentyl, isopentyl, 1-methylpentyl, 1,3-dimethylbutyl, n-hexyl, 1-methylhexyl, n-heptyl, isoheptyl, 1,1,3,3-tetramethylbutyl, 1-methylheptyl, 3-methylheptyl, n-octyl, 2-ethylhexyl, 1,1,3-trimethylhexyl,

25 1,1,3,3-tetramethylpentyl, nonyl, decyl, undecyl, 1-methylundecyl, dodecyl, 1,1,3,3,5,5-hexamethylhexyl, tridecyl, tetradecyl, pentadecyl, hexadecyl, heptadecyl, or octadecyl.

C₁-C₁₈Alkoxy radicals are straight-chain or branched alkoxy radicals, e.g. methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, sec-butoxy, tert-butoxy, amyloxy, isoamyloxy or tert-amyloxy, heptyloxy, octyloxy, isoocetylxy, nonyloxy, decyloxy, undecyloxy, dodecyloxy, tetradecyloxy,

30 pentadecyloxy, hexadecyloxy, heptadecyloxy and octadecyloxy.

C₂-C₁₈Alkenyl radicals are straight-chain or branched alkenyl radicals, such as e.g. vinyl, allyl, methallyl, isopropenyl, 2-butenyl, 3-butenyl, isobutenyl, n-penta-2,4-dienyl, 3-methylbut-2-enyl, n-oct-2-enyl, n-dodec-2-enyl, isododecanyl, n-dodec-2-enyl or n-octadec-4-enyl.

C₂₋₂₄Alkynyl is straight-chain or branched and preferably C₂₋₈alkynyl, which may be

35 unsubstituted or substituted, such as, for example, ethynyl, 1-propyn-3-yl, 1-butyn-4-yl, 1-pentyn-5-yl, 2-methyl-3-butyn-2-yl, 1,4-pentadiyn-3-yl, 1,3-pentadiyn-5-yl, 1-hexyn-6-yl,

cis-3-methyl-2-penten-4-yn-1-yl, *trans*-3-methyl-2-penten-4-yn-1-yl, 1,3-hexadiyn-5-yl, 1-octyn-8-yl, 1-nonyl-9-yl, 1-decyn-10-yl or 1-tetracosyn-24-yl, C₄-C₁₈cycloalkyl is preferably C₅-C₁₂cycloalkyl, such as, for example, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclononyl, cyclodecyl, cyclododecyl. Cyclohexyl and cyclododecyl

5 are most preferred.

The term "aryl group" is typically C₆-C₃₀aryl, such as phenyl, indenyl, azulenyl, naphthyl, biphenyl, terphenyl or quadphenyl, as-indacenyl, s-indacenyl, acenaphthyl, phenanthryl, fluoranthenyl, triphenyl, chrysenyl, naphthacen, picenyl, perylenyl, pentaphenyl, hexacenyl, pyrenyl, anthracenyl, terphenyl or quadphenyl, which can be

10 substituted or unsubstituted, preferably phenyl, 1-naphthyl, 2-naphthyl, 9-phenanthryl, 2- or 9-fluorenyl, 3- or 4-biphenyl, which may be unsubstituted or substituted. Examples of C₆-C₁₈aryl are phenyl, 4-methylphenyl, 4-methoxyphenyl, 1-naphthyl, 2-naphthyl, 3- or 4-biphenyl, 9-phenanthryl, 2- or 9-fluorenyl, which may be unsubstituted or substituted.

C₇-C₂₄aralkyl radicals are preferably C₇-C₁₅aralkyl radicals, which may be substituted, such

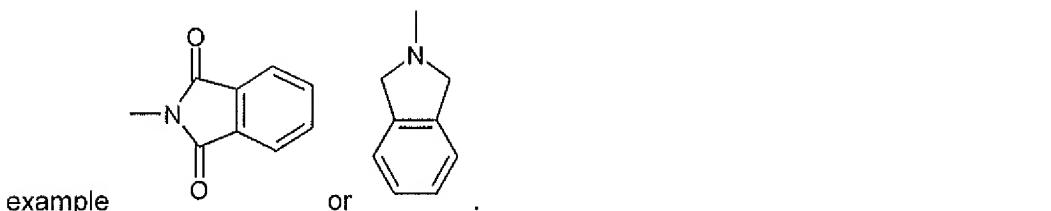
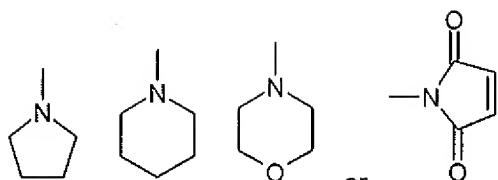
15 as, for example, benzyl, 2-benzyl-2-propyl, β -phenethyl, α,α -dimethylbenzyl, ω -phenyl-butyl, ω -phenyl-octyl, ω -phenyl-dodecyl or 3-methyl-5-(1',1',3',3'-tetramethyl-butyl)-benzyl,

The term "heteroaryl group", especially C₂-C₃₀heteroaryl, is a ring, wherein nitrogen, oxygen or sulfur are the possible hetero atoms, and is typically an unsaturated heterocyclic radical with five to 18 atoms having at least six conjugated π -electrons such as thienyl,

20 benzo[b]thienyl, dibenzo[b,d]thienyl, thianthrenyl, furyl, furfuryl, 2H-pyranyl, benzofuranyl, isobenzofuranyl, 2H-chromenyl, xanthenyl, dibenzofuranyl, phenoxythienyl, pyrrolyl, imidazolyl, pyrazolyl, pyridyl, bipyridyl, triazinyl, pyrimidinyl, pyrazinyl, 1H-pyrrolizinyl, isoindolyl, pyridazinyl, indolizinyl, isoindolyl, indolyl, 3H-indolyl, phthalazinyl, naphthyridinyl, quinoxalinyl, quinazolinyl, cinnolinyl, indazolyl, purinyl, quinolizinyl, chinolyl, isoquinolyl,

25 phthalazinyl, naphthyridinyl, chinoxalinyl, chinazolinyl, cinnolinyl, pteridinyl, carbazolyl, 4aH-carbazolyl, carbolinyl, benzotriazolyl, benzoxazolyl, phenanthridinyl, acridinyl, perimidinyl, phenanthrolinyl, phenazinyl, isothiazolyl, phenothiazinyl, isoxazolyl, furazanyl or phenoxazinyl, preferably the above-mentioned mono- or bicyclic heterocyclic radicals, which may be unsubstituted or substituted.

30 Examples of a five or six membered ring formed by R⁵ and R⁶ and R¹⁰³ and R¹⁰⁴, respectively are heterocycloalkanes or heterocycloalkenes having from 3 to 5 carbon atoms which can have one additional hetero atom selected from nitrogen, oxygen and sulfur, for example



Possible substituents of the above-mentioned groups are C₁-C₈alkyl, a hydroxyl group, a mercapto group, C₁-C₈alkoxy, C₁-C₈alkylthio, halogen, halo-C₁-C₈alkyl, a cyano group, an aldehyde group, a ketone group, a carboxyl group, an ester group, a carbamoyl group, an amino group, a nitro group or a silyl group.

As described above, the aforementioned radicals may be substituted by E and/or, if desired, 10 interrupted by D. Interruptions are of course possible only in the case of radicals containing at least 2 carbon atoms connected to one another by single bonds; C₆-C₁₈aryl is not interrupted; interrupted arylalkyl or alkylaryl contains the unit D in the alkyl moiety. C₁-C₁₈alkyl substituted by one or more E and/or interrupted by one or more units D is, for example, (CH₂CH₂O)_n-R^X, where n is a number from the range 1-9 and R^X is H or C₁-C₁₀alkyl 15 or C₂-C₁₀alkanoyl (e.g. CO-CH(C₂H₅)C₄H₉), CH₂-CH(OR^Y)-CH₂-O-R^Y, where R^Y is C₁-C₁₈alkyl, C₅-C₁₂cycloalkyl, phenyl, C₇-C₁₅phenylalkyl, and R^Y embraces the same definitions as R^Y or is H; C₁-C₈alkylene-COO-R^Z, e.g. CH₂COOR_Z, CH(CH₃)COOR^Z, C(CH₃)₂COOR^Z, where R^Z is H, C₁-C₁₈alkyl, (CH₂CH₂O)₁₋₉-R^X, and R^X embraces the definitions indicated above; 20 CH₂CH₂-O-CO-CH=CH₂; CH₂CH(OH)CH₂-O-CO-C(CH₃)=CH₂.

To obtain organic layers of this invention with the proper T_g, or glass transition temperature, it is advantageous that the present organic compounds have a melting point greater than about 150°C, for example greater than about 200°C, for example greater than about 250°C, for instance greater than about 300°C.

The electroluminescent devices of the present invention are otherwise designed as is known in the art, for example as described in U.S. Pat. Nos. 5,518,824, 6,225,467, 6,280,859, 5,629,389, 5,486,406, 5,104,740, 5,116,708 and 6,057,048, the relevant disclosures of which are hereby incorporated by reference.

5 For example, organic EL devices contain one or more layers such as: substrate; base electrode; hole-injecting layer; hole transporting layer; emitter layer; electron-transporting layer; electron-injecting layer; top electrode; contacts and encapsulation. This structure is a general case and may have additional layers or may be simplified by omitting layers so that one layer performs a plurality of tasks. For instance, the simplest 10 organic EL device consists of two electrodes which sandwich an organic layer that performs all functions, including the function of light emission.

A preferred EL device comprises in this order:

(a) an anode,
15 (b) a hole injecting layer and/or a hole transporting layer,
(c) a light-emitting layer,
(d) optionally an electron transporting layer and
(e) a cathode.

20 In particular, the present organic compounds function as light emitters and are contained in the light emission layer or form the light-emitting layer. The light emitting compounds of this invention exhibit intense fluorescence in the solid state and have excellent electric-field-applied light emission characteristics. Further, the light emitting compounds of this invention are excellent in the injection of holes from a metal electrode and the transportation of holes; as well as being excellent in the injection of 25 electrons from a metal electrode and the transportation of electrons. They are effectively used as light emitting materials and may be used in combination with other hole transporting materials, other electron transporting materials or other dopants. The organic compounds of the present invention form uniform thin films. The light emitting 30 layers may therefore be formed of the present organic compounds alone. Alternatively, the light-emitting layer may contain a known light-emitting material, a known dopant, a known hole transporting material or a known electron transporting material as required. In the organic EL device, a decrease in the brightness and life caused by quenching can be prevented by forming it as a multi-layered structure. The light-emitting 35 material, a dopant, a hole-injecting material and an electron-injecting material may be used in combination as required. Further, a dopant can improve the light emission brightness and the

light emission efficiency, and can attain the red or blue light emission. Further, each of the hole transporting zone, the light-emitting layer and the electron transporting zone may have the layer structure of at least two layers. In the hole transporting zone in this case, a layer to which holes are injected from an electrode is called "hole-injecting layer", and a layer which receives holes from the hole-injecting layer and transports the holes to a light-emitting layer is called "hole transporting layer". In the electron transporting zone, a layer to which electrons are injected from an electrode is called "electron-injecting layer", and a layer which receives electrons from the electron-injecting layer and transports the electrons to a light-emitting layer is called "electron transporting layer". These layers are selected and used depending upon factors such as the energy level and heat resistance of materials and adhesion to an organic layer or metal electrode.

The light-emitting material or the dopant which may be used in the light-emitting layer together with the organic compounds of the present invention includes for example anthracene, naphthalene, phenanthrene, pyrene, tetracene, coronene, chrysene, fluorescein, 15 perylene, phthaloperylene, naphthaloperylene, perinone, phthaoperinone, naphthaloperinone, diphenylbutadiene, tetraphenylbutadiene, coumarine, oxadiazole, aldazine, bisbenzoxazoline, bisstyryl, pyrazine, cyclopentadiene, quinoline metal complex, aminoquinoline metal complex, benzoquinoline metal complex, imine, diphenylethylene, vinyl anthracene, diaminocarbazole, pyran, thiopyran, polymethine, merocyanine, an imidazole-20 chelated oxynoid compound, quinacridone, rubrene, and fluorescent dyestuffs for a dyestuff laser or for brightening.

If the pyrimidine compounds of the formula I are used as host, the weight ratio of the pyrimidine compound (host) to the guest compound is in general 50:50 to 99.99:0.01, preferably 90:10 to 99.99:0.01, more preferably 95:5 to 99.9:0.1, most preferably 98:2 to 25 99.9:0.1.

The pyrimidine compounds of the present invention and the above compound or compounds that can be used in a light-emitting layer may be used in any mixing ratio for forming a light-emitting layer. That is, the organic compounds of the present invention may provide a main component for forming a light-emitting layer, or they may be a doping material in another 30 main material, depending upon a combination of the above compounds with the organic compounds of the present invention.

The hole-injecting material is selected from compounds which are capable of transporting holes, are capable of receiving holes from the anode, have an excellent effect of injecting holes to a light-emitting layer or a light-emitting material, prevent the movement of excitons 35 generated in a light-emitting layer to an electron-injecting zone or an electron-injecting material and have the excellent capability of forming a thin film. Suitable hole-injecting

materials include for example a phthalocyanine derivative, a naphthalocyanine derivative, a porphyrin derivative, oxazole, oxadiazole, triazole, imidazole, imidazolone, imidazolthione, pyrazoline, pyrazolone, tetrahydroimidazole, oxazole, oxadiazole, hydrazone, acylhydrazone, polyarylalkane, stilbene, butadiene, benzidine type triphenylamine, styrylamine type triphenylamine, diamine type triphenylamine, derivatives of these, and polymer materials such as polyvinylcarbazole, polysilane and an electroconducting polymer.

In the organic EL device of the present invention, the hole-injecting material which is more effective is an aromatic tertiary amine derivative or a phthalocyanine derivative. Although not specially limited, specific examples of the tertiary amine derivative include triphenylamine,

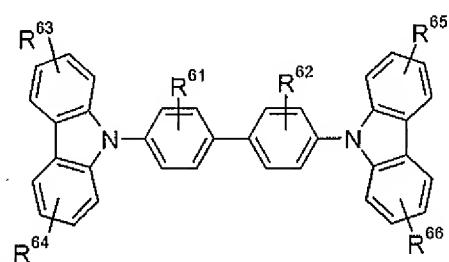
tritylamine, tolyldiphenylamine, N,N'-diphenyl-N,N'-(3-methylphenyl)-1,1-biphenyl-4,4'-diamine, N,N,N',N'-tetra(4-methylphenyl)-1,1'-phenyl-4,4'-diamine, N,N,N',N'-tetra(4-methylphenyl)-1,1'-biphenyl-4,4'-diamine, N,N'-diphenyl-N,N'-di(1-naphthyl)-1,1'-biphenyl-4,4'-diamine, N,N'-di(methylphenyl)-N,N'-di(4-n-butylphenyl)-phenanthrene-9,10-diamine, 4,4', 4"-tris(3-methylphenyl)-N-phenylamino)triphenylamine, 1,1-bis(4-di-p-

tolylaminophenyl)cyclohexane, and oligomers or polymers having aromatic tertiary amine structures of these.

Although not specially limited, specific examples of the phthalocyanine (Pc) derivative include phthalocyanine derivatives or naphthalocyanine derivatives such as H₂Pc, CuPc, CoPc, NiPc, ZnPc, PdPc, FePc, MnPc, ClAlPc, ClGaPc, ClInPc, ClSnPc, Cl₂SiPc, (HO)AlPc, (HO)GaPc, VOPc, TiOPc, MoOPc, and GaPc-O-GaPc.

The hole transporting layer can reduce the driving voltage of the device and improve the confinement of the injected charge recombination within the pyrimidin light emitting layer. Any conventional suitable aromatic amine hole transporting materials described for the hole-injecting layer may be selected for forming this layer.

A preferred class of hole transporting materials is comprised of 4,4'-bis(9-carbazolyl)-1,1'-biphenyl compounds of the formula



wherein R⁶¹ and R⁶² is a hydrogen atom or an C₁-C₃alkyl group; R⁶³ through R⁶⁶ are substituents independently selected from the group consisting of hydrogen, a C₁-C₆alkyl group, a C₁-C₆alkoxy group, a halogen atom, a dialkylamino group, a C₆-C₃₀aryl group, and

the like. Illustrative examples of 4,4'-bis(9-carbazolyl)-1,1'-biphenyl compounds include 4,4'-bis(9-carbazolyl)-1,1'-biphenyl and 4,4'-bis(3-methyl-9-carbazolyl)-1,1'-biphenyl, and the like.

The electron transporting layer is not necessarily required for the present device, but is optionally and preferably used for the primary purpose of improving the electron injection characteristics of the EL devices and the emission uniformity. Illustrative examples of electron transporting compounds, which can be utilized in this layer, include the metal chelates of 8-hydroxyquinoline as disclosed in U.S. Pat. Nos. 4,539,507, 5,151,629, and 5,150,006, the disclosures of which are totally incorporated herein by reference.

Although not specially limited, specific examples of the metal complex compound include

lithium 8-hydroxyquinolinate, zinc bis(8-hydroxyquinolinate), copper bis(8-

hydroxyquinolinate), manganese bis(8-hydroxyquinolinate), aluminum tris(8-

hydroxyquinolinate), aluminum tris(2-methyl-8-hydroxyquinolinate), gallium tris(8-

hydroxyquinolinate), beryllium bis(10-hydroxybenzo[h]quinolinate), zinc bis(10-

hydroxybenzo[h]quinolinate), chlorogallium bis(2-methyl-8-quinolinate), gallium bis(2-methyl-

8-quinolinate)(o-cresolate), aluminum bis(2-methyl-8-quinolinate)(1-naphtholate), gallium bis(2-methyl-8-quinolinate)(2-naphtholate), gallium bis(2-methyl-8-quinolinate)phenolate, zinc

bis(o-(2-benzoxazolyl)phenolate), zinc bis(o-(2-benzothiazolyl)phenolate) and zinc bis(o-(2-benzotriazolyl)phenolate). The nitrogen-containing five-membered derivative is preferably an

oxazole, thiazole, thiadiazole, or triazole derivative. Although not specially limited, specific examples of the above nitrogen-containing five-membered derivative include 2,5-bis(1-

phenyl)-1,3,4-oxazole, 1,4-bis(2-(4-methyl-5-phenyloxazolyl)benzene, 2,5-bis(1-phenyl)-

1,3,4-thiazole, 2,5-bis(1-phenyl)-1,3,4-oxadiazole, 2-(4'-tert-butylphenyl)-5-(4"-biphenyl)1,3,4-oxadiazole, 2,5-bis(1-naphthyl)-1,3,4-oxadiazole, 1,4-bis[2-(5-phenyloxadiazolyl)]benzene,

1,4-bis[2-(5-phenyloxadiazolyl)-4-tert-butylbenzene], 2-(4'-tert-butylphenyl)-5-(4"-biphenyl)-

1,3,4-thiadiazole, 2,5-bis(1-naphthyl)-1,3,4-thiadiazole, 1,4-bis[2-(5-phenylthiazolyl)]benzene, 2-(4'-tert-butylphenyl)-5-(4"-biphenyl)-1,3,4-triazole, 2,5-bis(1-naphthyl)-1,3,4-triazole and

1,4-bis[2-(5-phenyltriazolyl)]benzene. Another class of electron transport materials are oxadiazole metal chelates, such as bis[2-(2-hydroxyphenyl)-5-phenyl-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-phenyl-1,3,4-oxadiazolato]beryllium; bis[2-(2-

hydroxyphenyl)-5-(1-naphthyl)-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-(1-

naphthyl)-1,3,4-oxadiazolato]beryllium; bis[5-biphenyl-2-(2-hydroxyphenyl)-1,3,4-oxadiazolato]zinc; bis[5-biphenyl-2-(2-hydroxyphenyl)-1,3,4-oxadiazolato]beryllium; bis(2-

hydroxyphenyl)-5-phenyl-1,3,4-oxadiazolato]lithium; bis[2-(2-hydroxyphenyl)-5-p-tolyl-1,3,4-

oxadiazolato]zinc; bis 2-(2-hydroxyphenyl)-5-p-tolyl-1,3,4-oxadiazolato]beryllium; bis[5-(p-

tert-butylphenyl)-2-(2-hydroxyphenyl)-1,3,4-oxadiazolato]zinc; bis[5-(p-tert-butylphenyl)-2-(2-

hydroxyphenyl)-1,3,4-oxadiazolato]beryllium; bis[2-(2-hydroxyphenyl)-5-(3-fluorophenyl)-

1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-(4-fluorophenyl)-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-(4-fluorophenyl)-1,3,4-oxadiazolato]beryllium; bis[5-(4-chlorophenyl)-2-(2-hydroxyphenyl)-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-(4-methoxyphenyl)-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxy-4-methylphenyl)-5-phenyl-1,3,4-oxadiazolato]zinc; bis[2-.alpha.-(2-hydroxynaphthyl)-5-phenyl-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-p-pyridyl-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-p-pyridyl-1,3,4-oxadiazolato]beryllium; bis[2-(2-hydroxyphenyl)-5-(2-thiophenyl)-1,3,4-oxadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-phenyl-1,3,4-thiadiazolato]zinc; bis[2-(2-hydroxyphenyl)-5-phenyl-1,3,4-thiadiazolato]beryllium; bis[2-(2-hydroxyphenyl)-5-(1-naphthyl)-1,3,4-thiadiazolato]zinc; and bis[2-(2-hydroxyphenyl)-5-(1-naphthyl)-1,3,4-thiadiazolato]beryllium, and the like.

In the organic EL device of the present invention, the light-emitting layer may contain, in addition to the light-emitting organic material of the present invention, at least one of other light-emitting material, other dopant, other hole-injecting material and other electron-injecting material. For improving the organic EL device of the present invention in the stability against temperature, humidity and ambient atmosphere, a protective layer may be formed on the surface of the device, or the device as a whole may be sealed with a silicone oil, or the like.

The electrically conductive material used for the anode of the organic EL device is suitably selected from those materials having a work function of greater than 4 eV. The electrically conductive material includes carbon, aluminum, vanadium, iron, cobalt, nickel, tungsten, silver, gold, platinum, palladium, alloys of these, metal oxides such as tin oxide and indium oxide used for ITO substrates or NESA substrates, and organic electroconducting polymers such as polythiophene and polypyrrole.

The electrically conductive material used for the cathode is suitably selected from those having a work function of smaller than 4 eV. The electrically conductive material includes magnesium, calcium, tin, lead, titanium, yttrium, lithium, ruthenium, manganese, aluminum and alloys of these, while the electrically conductive material shall not be limited to these. Examples of the alloys include magnesium/silver, magnesium/indium and lithium/aluminum, while the alloys shall not be limited to these. Each of the anode and the cathode may have a layer structure formed of two layers or more as required.

For the effective light emission of the organic EL device, at least one of the electrodes is desirably sufficiently transparent in the light emission wavelength region of the device. Further, the substrate is desirably transparent as well. The transparent electrode is produced from the above electrically conductive material by a deposition method or a sputtering method such that a predetermined light transmittance is secured. The electrode on the light emission surface side has for instance a light transmittance of at least 10%. The

substrate is not specially limited so long as it has adequate mechanical and thermal strength and has transparency. For example, it is selected from glass substrates and substrates of transparent resins such as a polyethylene substrate, a polyethylene terephthalate substrate, a polyether sulfone substrate and a polypropylene substrate.

5 In the organic EL device of the present invention, each layer can be formed by any one of dry film forming methods such as a vacuum deposition method, a sputtering method, a plasma method and an ion plating method and wet film forming methods such as a spin coating method, a dipping method and a flow coating method. The thickness of each layer is not specially limited, while each layer is required to have a proper thickness. When the layer
10 thickness is too large, inefficiently, a high voltage is required to achieve predetermined emission of light. When the layer thickness is too small, the layer is liable to have a pinhole, etc., so that sufficient light emission brightness is hard to obtain when an electric field is applied. The thickness of each layer is for example in the range of from about 5 nm to about 10 μ m, for instance about 10 nm to about 0.2 μ m.

15 In the wet film forming method, a material for forming an intended layer is dissolved or dispersed in a proper solvent such as ethanol, chloroform, tetrahydrofuran and dioxane, and a thin film is formed from the solution or dispersion. The solvent shall not be limited to the above solvents. For improving the film formability and preventing the occurrence of pinholes in any layer, the above solution or dispersion for forming the layer may contain a proper resin
20 and a proper additive. The resin that can be used includes insulating resins such as polystyrene, polycarbonate, polyarylate, polyester, polyamide, polyurethane, polysulfone, polymethyl methacrylate, polymethyl acrylate and cellulose, copolymers of these, photoconductive resins such as poly-N-vinylcarbozole and polysilane, and electroconducting polymers such as polythiophene and polypyrrole. The above additive includes an antioxidant,
25 an ultraviolet absorbent and a plasticizer.

When the light-emitting organic material of the present invention is used in a light-emitting layer of an organic EL device, an organic EL device can be improved in organic EL device characteristics such as light emission efficiency and maximum light emission brightness. Further, the organic EL device of the present invention is remarkably stable against heat and
30 electric current and gives a usable light emission brightness at a low actuation voltage. The problematic deterioration of conventional devices can be remarkably decreased.

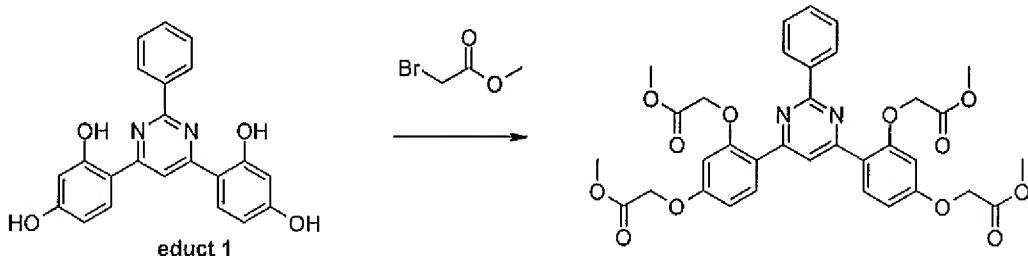
The organic EL device of the present invention has significant industrial values since it can be adapted for a flat panel display of an on-wall television set, a flat light-emitting device, a light source for a copying machine or a printer, a light source for a liquid crystal display or
35 counter, a display signboard and a signal light.

The material of the present invention can be used in the fields of an organic EL device, an electrophotographic photoreceptor, a photoelectric converter, a solar cell, an image sensor, dye lasers and the like.

5 The following Examples illustrate the invention. In the Examples and throughout this application, the term light emitting material means the present pyrimidine compounds.

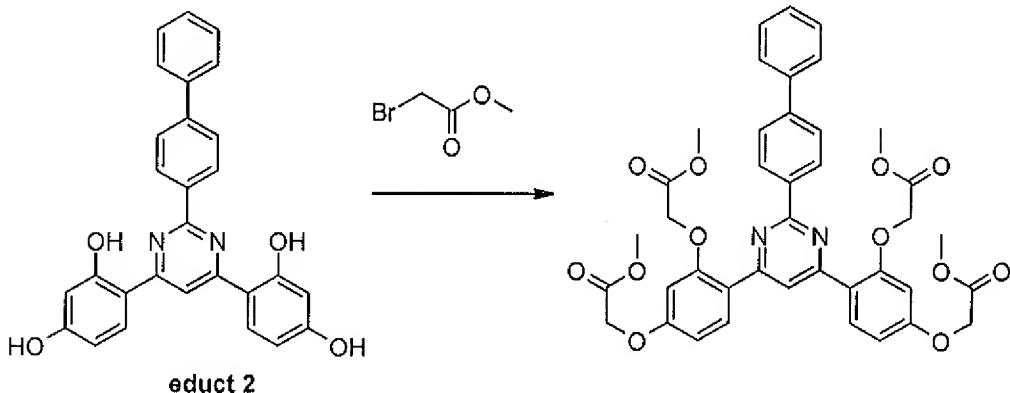
Examples

Example 1 (A-1)



To 1.00 g (2.69 mmol) of educt 1 (prepared according to US-A-3,442,898 from resorcinol and 4,6-dichloro-2-phenyl-pyrimidine; J. Org. Chem. 1988, 53, 4137) in 20 ml water free DMF 3.29 g (21.5 mmol) methyl-bromoacetate and 2.97 g (21.5 mmol) potassium carbonate are added. The reaction mixture is stirred at 100 °C under nitrogen for 2 h. The reaction mixture 15 is diluted with water. The organic phase is extracted with dichloromethane und dried with magnesium sulfate. The solvent is removed. After chromatography of the crude product on silica gel with toluene / ethyl acetate 8/2 the desired product is obtained. Melting point: 178.0 – 179.0 °C.

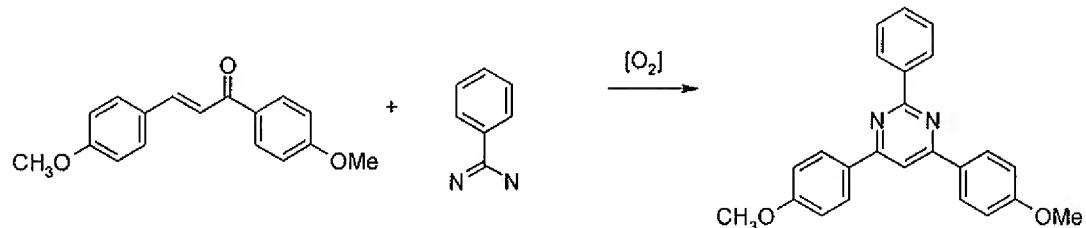
20 Example 2 (A-2)



To 520 mg (1.16 mmol) of educt 2 (prepared according to US-A-3,442,898 from resorcinol and 4,6-dichloro-2-biphenyl-pyrimidine; EP-A-96657) in 20 ml water free DMF 1.42 g (9.28

mmol) methyl-bromoacetate and 1.28 g (9.28 mmol) potassium carbonate are added. The reaction mixture is stirred at 100 °C under nitrogen for 2 h. The reaction mixture is diluted with water. The organic phase is extracted with dichloromethane und dried with magnesium sulfate. The solvent is removed. After chromatography of the crude product on silica gel with toluene / ethyl acetate 9/1 and later 7/3 the desired product is obtained. Melting point: 119.5 – 121.5 °C.

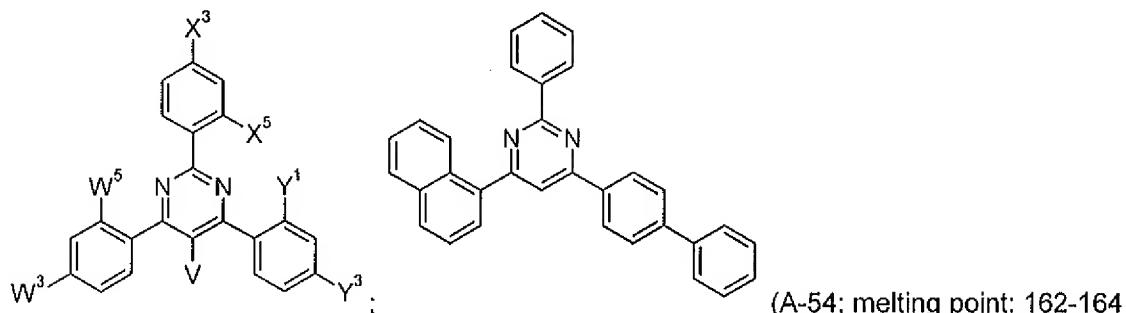
Example 3 (A-3)



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To 4.31 g (16.1 mmol) of 1,3-bis(4-methoxyphenyl)-2-propen-1-one in 25 ml water free ethanol 1.89 g (8.11 mmol) biphenylbenzimidine hydrochlorid are added. A solution of 1.07 g (19.08 mmol) potassium hydroxide is added during 15 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 24 h. The reaction mixture is poured 15 into water. The solid is filtered off and washed with water. The product is crystallized 2 times from acetic acid (96-98 %). Melting point: 168-169 °C.

Compounds A-4 to A-54 can be obtained in a manner analogous to Examples 1 to 5.

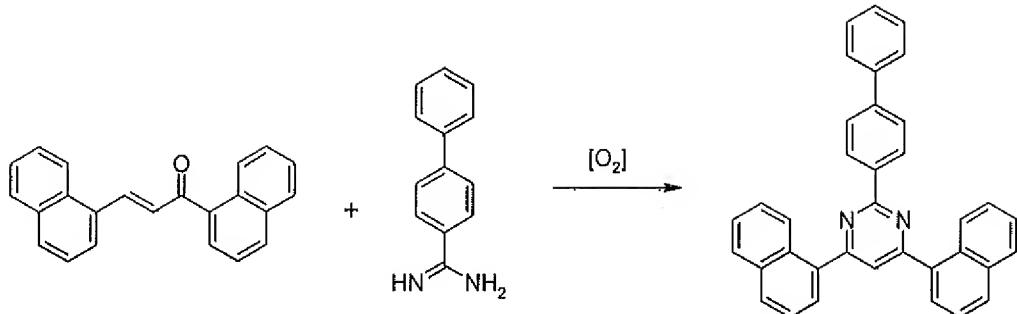


Cpd.	X ³	Y ³	W ³	X ⁵	Y ¹	W ⁶	V
A-1	H	OCH ₂ COOMe	OCH ₂ COOMe	H	OCH ₂ COOMe	OCH ₂ COOMe	H
A-2	Ph	OCH ₂ COOMe	OCH ₂ COOMe	H	OCH ₂ COOMe	OCH ₂ COOMe	H
A-3	H	OMe	OMe	H	H	H	H
A-4 ¹⁾	H	Ph	Ph	H	H	H	H
A-5	H	OMe	OMe	H	OMe	OMe	H
A-6	H	OBu	OBu	H	H	H	H
A-7	H	OBu	OBu	H	OBu	OBu	H
A-8	OMe	OMe	OMe	OMe	OMe	OMe	H
A-9	OMe	Br	Br	OMe	OMe	OMe	H
A-10	Ph	Br	Br	H	H	H	H
A-11	OMe	Ph	Ph	OMe	OMe	OMe	H
A-12	Ph	Ph	Ph	H	H	H	H
A-13	SMe	SMe	SMe	H	H	H	H
A-14	H	SMe	SMe	H	H	H	H
A-15	H	OCH ₂ CH ₂ OAc	OCH ₂ CH ₂ OAc	H	H	H	H
A-16	H	OCH ₂ CH ₂ OAc	OCH ₂ CH ₂ OAc	H	OCH ₂ CH ₂ OAc	OCH ₂ CH ₂ OAc	H
A-17	H	OCH ₂ CH ₂ OCH ₂ Et	OCH ₂ CH ₂ OCH ₂ Et	H	H	H	H
A-18	H	OCH ₂ CH ₂ OCH ₂ Et	OCH ₂ CH ₂ OCH ₂ Et	H	OCH ₂ CH ₂ OCH ₂ Et	OCH ₂ CH ₂ OCH ₂ Et	H
A-19	H	N(CH ₃) ₂	N(CH ₃) ₂	H	H	H	H
A-20	H	-N ⁺ 	-N ⁺ 	H	H	H	H
A-21	Ph	N(CH ₃) ₂	N(CH ₃) ₂	H	H	H	H
A-22	Ph	-N ⁺ 	-N ⁺ 	H	H	H	H
A-23	H	CH ₃	CH ₃	H	H	H	H
A-24	Ph	CH ₃	CH ₃	H	H	H	H
A-25	Ph	Ph	Ph	H	H	H	CH ₃
A-26	OMe	OMe	OMe	H	H	H	CH ₃
A-27	H	Ph	Ph	H	H	H	CH ₃

¹⁾ melting point: 244-245°C.

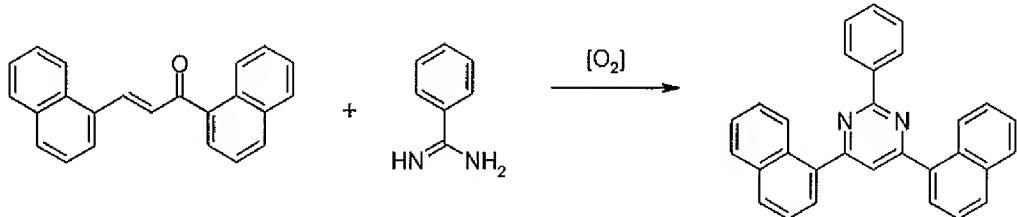
Cpd.	44		44		W ⁵	V
	X ³	Y ³	X ⁵	Y ¹		
A-28	H	OMe	OMe	H	H	CH ₃
A-29	H	OMe	OMe	H	OMe	CH ₃
A-30	H	OBu	OBu	H	H	OMe
A-31	H	OBu	OBu	H	OBu	CH ₃
A-32	OMe	OMe	OMe	OMe	OMe	CH ₃
A-33	OMe	Br	Br	OMe	OMe	CH ₃
A-34	Ph	Br	Br	H	H	CH ₃
A-35	OMe	Ph	Ph	OMe	H	CH ₃
A-36	H	OCH ₂ COOMe	OCH ₂ COOMe	H	OCH ₂ COOMe	CH ₃
A-37	SMe	SMe	SMe	H	H	CH ₃
A-38	H	SMe	SMe	H	H	CH ₃
A-39	H	OCH ₂ CH ₂ Oac	OCH ₂ CH ₂ Oac	H	H	CH ₃
A-40	H	OCH ₂ CH ₂ Oac	OCH ₂ CH ₂ Oac	H	OCH ₂ CH ₂ Oac	CH ₃
A-41	H	OCH ₂ CH ₂ OCH ₂ Et	OCH ₂ CH ₂ OCH ₂ Et	H	H	CH ₃
A-42	H	OCH ₂ CH ₂ OCH ₂ Et	OCH ₂ CH ₂ OCH ₂ Et	H	OCH ₂ CH ₂ OCH ₂ Et	CH ₃
A-43	OBu	Ph	Ph	OBu	H	CH ₃
A-44	H	N(CH ₃) ₂	N(CH ₃) ₂	H	H	CH ₃
A-45	H	-N ₂ +	-N ₂ +	H	H	CH ₃
A-46	Ph	N(CH ₃) ₂	N(CH ₃) ₂	H	H	CH ₃
A-47	Ph	-N ₂ +	-N ₂ +	H	H	CH ₃
A-48	Ph	CH ₃	CH ₃	H	H	CH ₃
A-49	Ph	CH ₃	CH ₃	H	H	CH ₃
A-50	OMe	OMe	OMe	H	H	H
A-51 ¹⁾	H	H	H	H	H	H
A-52 ²⁾	Ph	H	H	H	H	H

¹⁾ melting point: 188-189°C.
²⁾ melting point: 196-198°C

Example 4 (B-1)

To 5.00 g (16.1 mmol) of 1,3-bis- α -naphthyl-2-propen-1-one (1) in 25 ml water free ethanol 1.89 g (8.11 mmol) biphenylbenzamidine hydrochloride are added. A solution of 1.07 g (19.08 mmol) potassium hydroxide in 25 ml water free ethanol is added during 15 min. Dry air was bubbled through the reaction mixture. The reaction mixture is refluxed for 24 h.

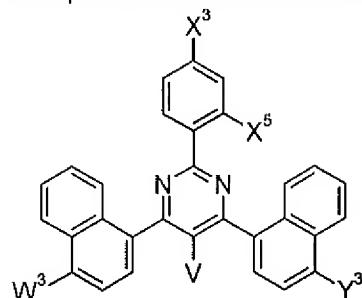
5 The reaction mixture is poured into water. The solid is filtered off and is washed with water. The product is crystallized 2 times from acetic acid (96-98 %). Melting point: 226-230 °C.

10 Example 5 (B-2)

To 5.00 g (16.1 mmol) of 1,3-bis- α -naphthyl-2-propen-1-one (1) in 25 ml water free ethanol 1.27 g (8.11 mmol) benzamidine hydrochloride are added. A solution of 1.07 g (19.08 mmol) potassium hydroxide in 25 ml water free ethanol is added during 15 min. Dry air is bubbled

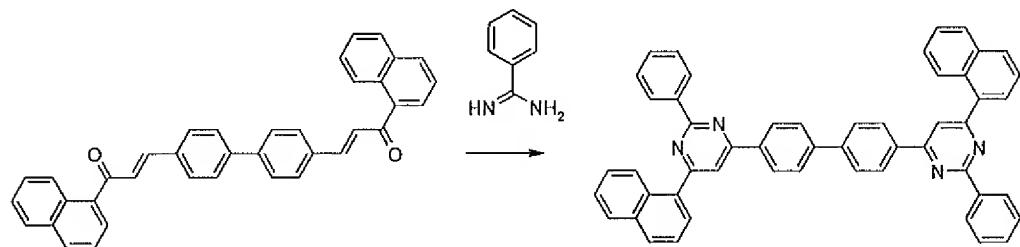
15 through the reaction mixture. The reaction mixture is refluxed for 24 h. The reaction mixture is poured into water. The solid is filtered off and is washed with water. The product is crystallized from acetic acid (96-98 %). Melting point: 179-180 °C.

Compounds B-3 to B-23 can be obtained in a manner analogous to Examples 4 and 5.



Cpd.	X ³	Y ³	W ³	X ⁵	V
B-1	Ph	H	H	H	H
B-2	H	H	H	H	H
B-3	OBu	H	H	OBu	H
B-4	OMe	H	H	OMe	H
B-5	OCH ₂ COOMe	H	H	OCH ₂ COOMe	H
B-6	H	OMe	OMe	H	H
B-7	OCH ₂ CH ₂ OCH ₂ Et	H	H	OCH ₂ CH ₂ OCH ₂ Et	H
B-8	OCH ₂ CH ₂ OAc	H	H	OCH ₂ CH ₂ OAc	H
B-9	OMe	OMe	OMe	OMe	H
B-10	OMe	OMe	OMe	OMe	H
B-11	H	CH ₃	CH ₃	H	H
B-12	Ph	CH ₃	CH ₃	H	H
B-13	Ph	H	H	H	CH ₃
B-14	OMe	H	H	OMe	CH ₃
B-15	OBu	H	H	OBu	CH ₃
B-16	H	H	H	H	CH ₃
B-17	OCH ₂ COOMe	H	H	OCH ₂ COOMe	CH ₃
B-18	H	OMe	OMe	H	CH ₃
B-19	OCH ₂ CH ₂ OCH ₂ Et	H	H	OCH ₂ CH ₂ OCH ₂ Et	CH ₃
B-20	OCH ₂ CH ₂ OAc	H	H	OCH ₂ CH ₂ OAc	CH ₃
B-21	H	CH ₃	CH ₃	H	CH ₃
B-22	Ph	CH ₃	CH ₃	H	CH ₃

5 Example 6 (C-1)



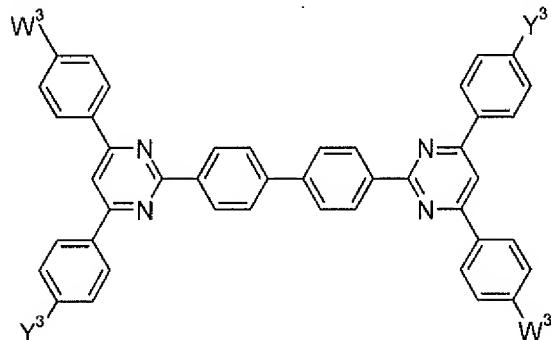
To 4.50 g (8.74 mmol) of the 2-propen-1-one derivative shown above in 50 ml water free ethanol 1.37 g (8.74 mmol) benzimidazole hydrochloride are added. A solution of 1.15 g (20.6 mmol) potassium hydroxide in 50 ml water free ethanol is added during 15 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 24 h, poured into

10

water and the water phase is extracted with dichloromethane. The organic phase is dried with magnesium sulfate, the solvent is removed by distillation and the remaining residue is purified by column chromatography (toluene / hexane 2/1).

5 $^1\text{H-NMR}$ (300 MHz, CDCl_3): δ = 8.75-8.72 (m, 4H); 8.45-8.37 (m, 12H); 8.04-7.82 (m, 6H); 7.67-7.50 (m, 12H).

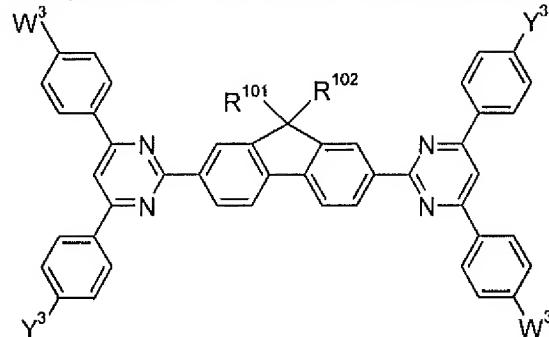
Compounds C-2 to C-24 can be obtained in a manner analogous to Example 6.



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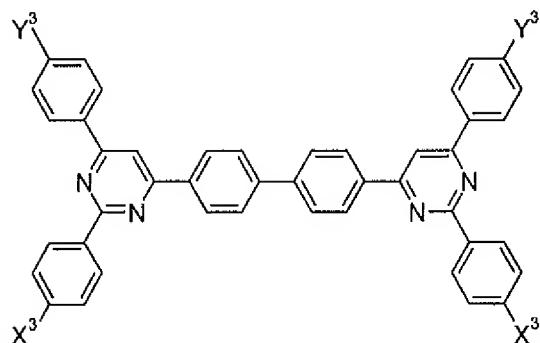
Cpd.	W^3	Y^3
C-2	H	H
C-3	H	H
C-4	H	H
C-5	H	H
C-6	H	H
C-7	OCH_3	OCH_3
C-8	OCH_3	OCH_3
C-9	OCH_3	OCH_3
C-10	OCH_3	OCH_3
C-11	OCH_3	OCH_3
C-12	SCH_3	SCH_3
C-13	SCH_3	SCH_3
C-14	SCH_3	SCH_3
C-15	SCH_3	SCH_3
C-16	SCH_3	SCH_3
C-17	SCH_3	SCH_3
C-18	$\text{N}(\text{CH}_3)_2$	$\text{N}(\text{CH}_3)_2$
C-19	$\text{N}(\text{CH}_3)_2$	$\text{N}(\text{CH}_3)_2$
C-20	$\text{N}(\text{CH}_3)_2$	$\text{N}(\text{CH}_3)_2$
C-21	$\text{N}(\text{CH}_3)_2$	$\text{N}(\text{CH}_3)_2$
C-22	$\text{N}(\text{CH}_3)_2$	$\text{N}(\text{CH}_3)_2$
C-23	$\text{N}(\text{CH}_3)_2$	$\text{N}(\text{CH}_3)_2$
C-24	$\text{N}(\text{CH}_3)_2$	$\text{N}(\text{CH}_3)_2$

Compounds D-1 to D-23 can be obtained in a manner analogous to Example 6.



Cpd.	R^{101}	R^{102}	W^3	Y^3
D-1	C_6H_{13}	C_6H_{13}	H	H
D-2	Bu	Bu	H	H
D-3	Et	Et	H	H
D-4	H	H	H	H
D-5	Ph	Ph	H	H
D-6	C_6H_{13}	C_6H_{13}	OCH_3	OCH_3
D-7	Bu	Bu	OCH_3	OCH_3
D-8	Et	Et	OCH_3	OCH_3
D-9	H	H	OCH_3	OCH_3
D-10	Ph	Ph	OCH_3	OCH_3
D-11	C_6H_{13}	C_6H_{13}	SCH_3	SCH_3
D-12	Bu	Bu	SCH_3	SCH_3
D-13	Et	Et	SCH_3	SCH_3
D-14	H	H	SCH_3	SCH_3
D-15	Ph	Ph	SCH_3	SCH_3
D-16	C_6H_{13}	C_6H_{13}	SCH_3	SCH_3
D-17	C_6H_{13}	C_6H_{13}	$N(CH_3)_2$	$N(CH_3)_2$
D-18	Bu	Bu	$N(CH_3)_2$	$N(CH_3)_2$
D-19	Et	Et	$N(CH_3)_2$	$N(CH_3)_2$
D-20	H	H	$N(CH_3)_2$	$N(CH_3)_2$
D-21	Ph	Ph	$N(CH_3)_2$	$N(CH_3)_2$
D-22	C_6H_{13}	C_6H_{13}	$N(CH_3)_2$	$N(CH_3)_2$
D-23	Bu	Bu	$N(CH_3)_2$	$N(CH_3)_2$

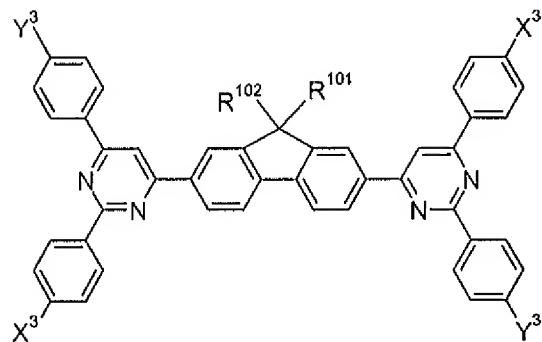
Compounds E-1 to E-35 can be obtained in a manner analogous to Example 6.



5

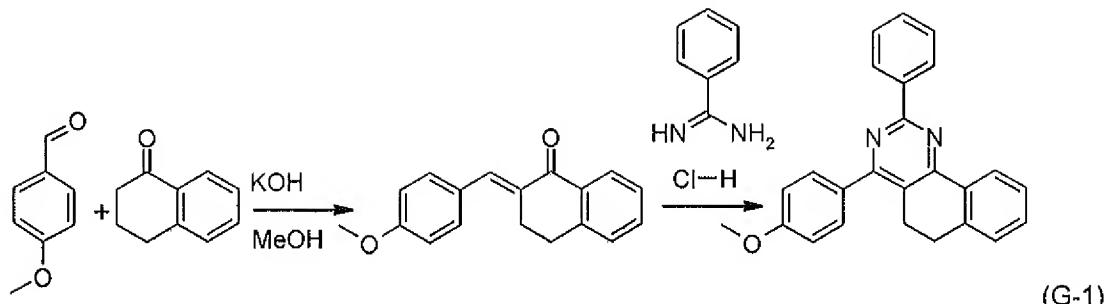
Cpd.	Y ³	X ³
E-1	H	H
E-2	H	H
E-3	H	H
E-4	H	H
E-5	H	H
E-6	OCH ₃	OCH ₃
E-7	OCH ₃	OCH ₃
E-8	OCH ₃	OCH ₃
E-9	OCH ₃	OCH ₃
E-10	OCH ₃	OCH ₃
E-11	OCH ₃	H
E-12	OCH ₃	H
E-13	OCH ₃	H
E-14	OCH ₃	H
E-15	OCH ₃	H
E-16	SCH ₃	SCH ₃
E-17	SCH ₃	SCH ₃
E-18	SCH ₃	SCH ₃
E-19	SCH ₃	SCH ₃
E-20	SCH ₃	SCH ₃
E-21	SCH ₃	H
E-22	SCH ₃	H
E-23	SCH ₃	H
E-24	SCH ₃	H
E-25	SCH ₃	H
E-26	N(CH ₃) ₂	N(CH ₃) ₂
E-27	N(CH ₃) ₂	N(CH ₃) ₂
E-28	N(CH ₃) ₂	N(CH ₃) ₂
E-29	N(CH ₃) ₂	N(CH ₃) ₂
E-30	N(CH ₃) ₂	N(CH ₃) ₂
E-31	N(CH ₃) ₂	H
E-32	N(CH ₃) ₂	H
E-33	N(CH ₃) ₂	H
E-34	N(CH ₃) ₂	H
E-35	N(CH ₃) ₂	H

Compounds F-1 to F-35 can be obtained in a manner analogous to Example 6.



Cpd.	R ¹⁰¹	R ¹⁰²	Y ³	X ³
F-1	C ₆ H ₁₃	C ₆ H ₁₃	H	H
F-2	Bu	Bu	H	H
F-3	Et	Et	H	H
F-4	H	H	H	H
F-5	Ph	Ph	H	H
F-6	C ₆ H ₁₃	C ₆ H ₁₃	OCH ₃	OCH ₃
F-7	Bu	Bu	OCH ₃	OCH ₃
F-8	Et	Et	OCH ₃	OCH ₃
F-9	H	H	OCH ₃	OCH ₃
F-10	Ph	Ph	OCH ₃	OCH ₃
F-11	C ₆ H ₁₃	C ₆ H ₁₃	OCH ₃	H
F-12	Bu	Bu	OCH ₃	H
F-13	Et	Et	OCH ₃	H
F-14	H	H	OCH ₃	H
F-15	Ph	Ph	OCH ₃	H
F-16	C ₆ H ₁₃	C ₆ H ₁₃	SCH ₃	SCH ₃
F-17	Bu	Bu	SCH ₃	SCH ₃
F-18	Et	Et	SCH ₃	SCH ₃
F-19	H	H	SCH ₃	SCH ₃
F-20	Ph	Ph	SCH ₃	SCH ₃
F-21	C ₆ H ₁₃	C ₆ H ₁₃	SCH ₃	H
F-22	Bu	Bu	SCH ₃	H
F-23	Et	Et	SCH ₃	H
F-24	H	H	SCH ₃	H
F-25	Ph	Ph	SCH ₃	H
F-26	C ₆ H ₁₃	C ₆ H ₁₃	N(CH ₃) ₂	N(CH ₃) ₂
F-27	Bu	Bu	N(CH ₃) ₂	N(CH ₃) ₂
F-28	Et	Et	N(CH ₃) ₂	N(CH ₃) ₂
F-29	H	H	N(CH ₃) ₂	N(CH ₃) ₂
F-30	Ph	Ph	N(CH ₃) ₂	N(CH ₃) ₂
F-31	C ₆ H ₁₃	C ₆ H ₁₃	N(CH ₃) ₂	H
F-32	Bu	Bu	N(CH ₃) ₂	H
F-33	Et	Et	N(CH ₃) ₂	H
F-34	H	H	N(CH ₃) ₂	H
F-35	Ph	Ph	N(CH ₃) ₂	H

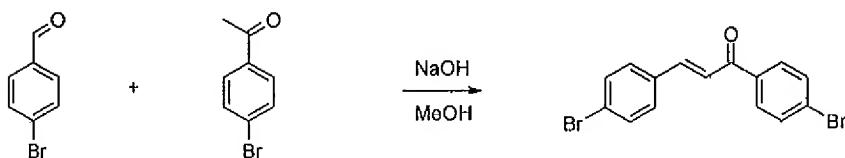
Example 7 (G-1)



To 14.6 g (0.100 mol) of tetralone and 15.0 g (1.10 mol) p-methoxy-benzaldehyd in 100 ml absolute methanol 660 mg potassium hydroxide are added. The reaction mixture is refluxed

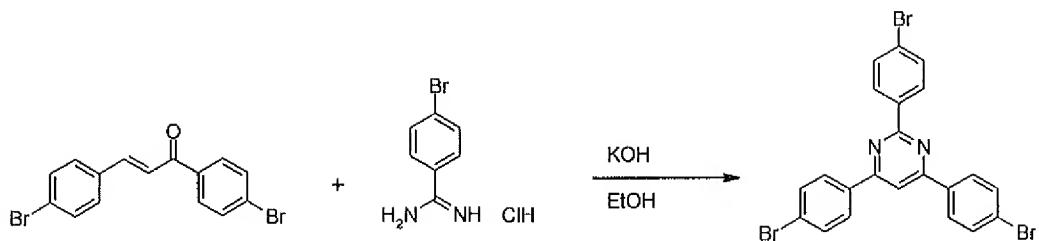
5 for 18 h under argon and then cooled to 25 °C. The formed product is filtered off and washed with methanol. To 6.61 g (25.0 mmol) of said product in 50 ml water free ethanol 1.96 g (12.5 mmol) benzamidine hydrochloride are added. A solution of 1.65 g (25.0 mmol) potassium hydroxide in 50 ml water free ethanol is added during 15 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 24 h and then poured into water. The
10 water phase is extracted with dichloromethane. The organic phase is dried with magnesium sulfate, the solvent is removed by distillation and the remaining residue is purified by column chromatography (toluene / hexane 1/1). The product G-1 having a melting point of 169 °C is obtained.

15 Example 8 (H-1)

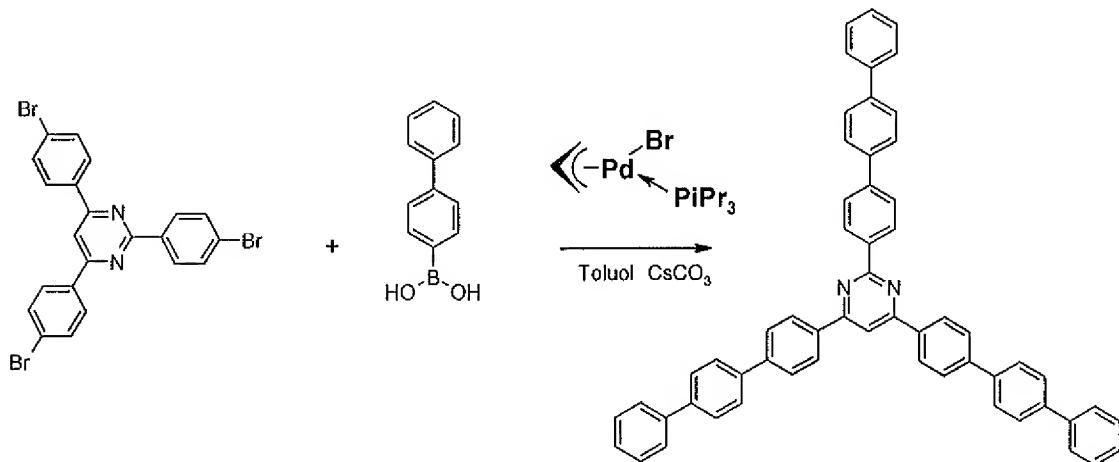


a) 24.9 g (0.134 mol) 4-bromo-benzaldehyde and 26.8 g 4-bromo-acetophenone (0.134 mol)

20 are dissolved under argon in 120 ml methanol. To this solution 0.270 g sodium hydroxide are added. The reaction mixture is stirred for 5 h. The formed yellow product is filtered off and is washed with water and then methanol. The product is dried in a vacuum oven (yield: 44.2 g (90 %), melting point: 183 – 184 °C).

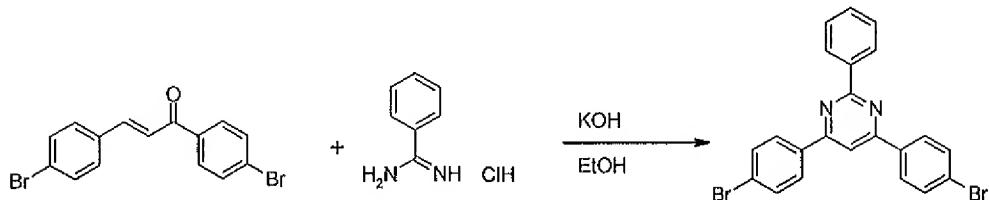


b) To 5.00 g (13.7 mmol) of 1,3-bis-(4-bromophenyl)-2-propen-1-one (1) in 25 ml water free ethanol 1.61 g (6.83 mmol) 4-bromo-benzamidine hydrochloride are added under argon. A solution of 0.90 g (16.1 mmol) potassium hydroxide in 25 ml water free ethanol is added 5 during 15 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 24 h and then poured into water. The product is filtered off, washed with ethanol and dried in a vacuum oven (melting point: 321 – 322 °C).

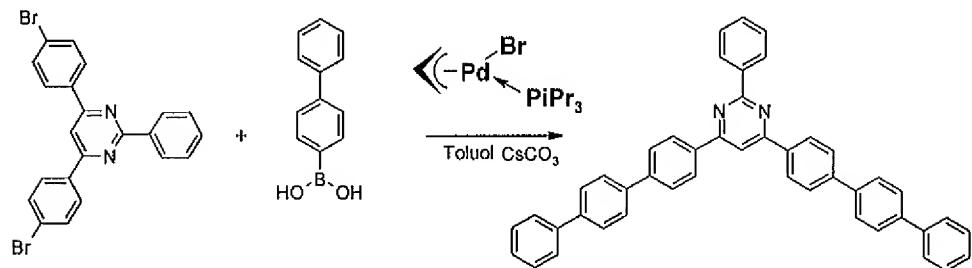


c) To 1.00 g (1.83 mmol) of tris-2,4,6-(4-bromo-phenyl)-pyrimidine and 1.27 g (6.42 mmol) of 4-biphenylboronic acid in 20 ml toluene 5.27 g (16.19 mmol) CsCO_3 in 3 ml water are added under argon. Approximately 10 mg of the Pd catalyst (WO 99/47474) are added and then the reaction mixture is refluxed for 5 h. The product is filtered off, washed with water and acetone. The product is dissolved in dichloromethane and filtered on silica gel. The solvent is 15 removed in vacuum. The obtained product has a melting point of 364 – 367 °C.

Example 9 (H-2)



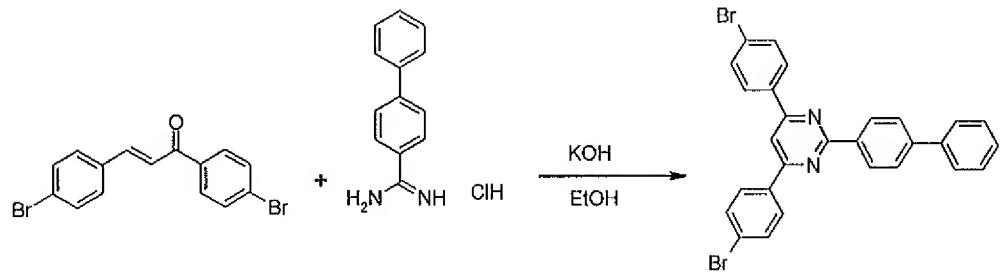
a) The product (melting point: 204 – 205 °C) is prepared according to the procedure given in example 8b).



b) The product (melting point: 343-345 °C) is prepared according to the procedure given in example 8c).

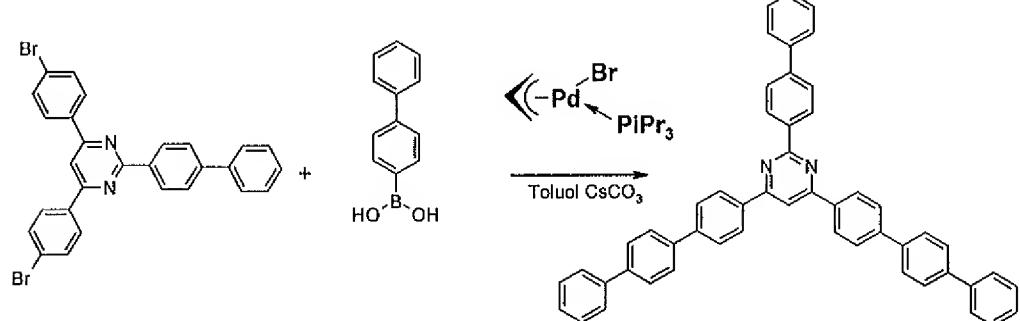
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Example 10 (H-3)



a) The product (melting point: 207 – 208 °C) is prepared according to the procedure given in example 8b).

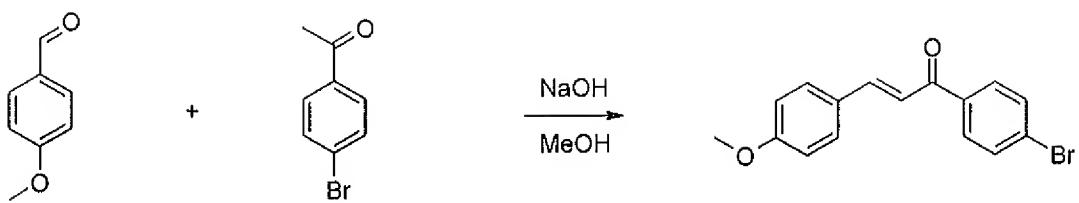
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b) The product (melting point: 364 – 367 °C) is prepared according to the procedure given in example 8c).

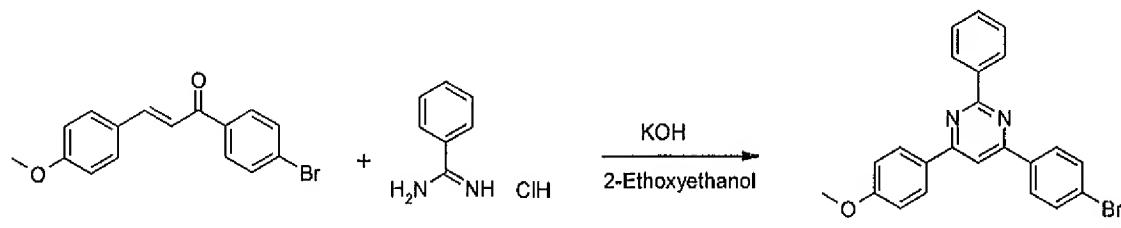
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Example 11 (H-4)

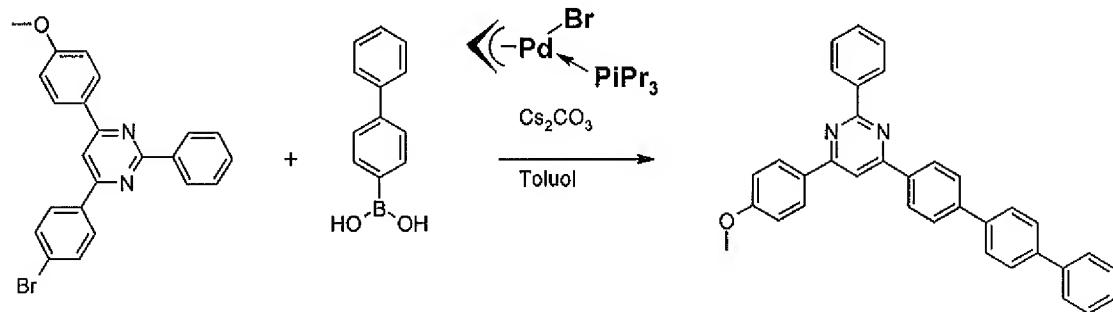


a) The product (melting point: 145-146 °C) is prepared according to the procedure given in example 8b) (yield: 86 %).

5

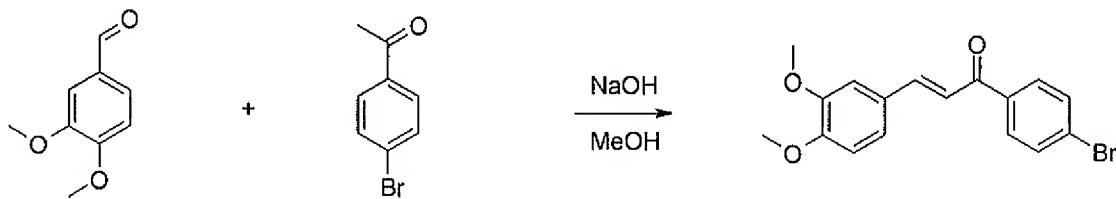


b) The product (melting point: 160 °C) is prepared according to the procedure given in example 8b) (yield: 72 %).



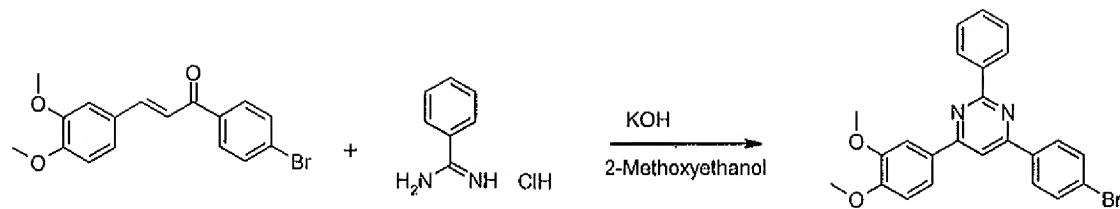
c) The product (melting point: 213-214 °C) is prepared according to the procedure given in example 8c) (yield: 53 %).

Example 12 (H-5)

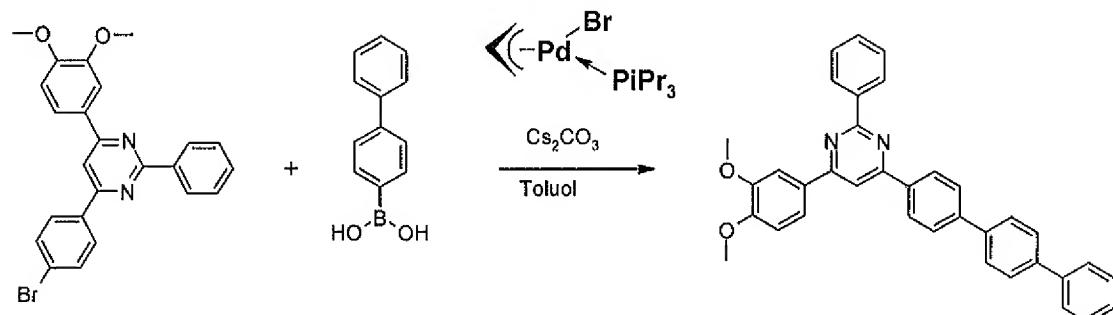


a) The product (melting point: 122-123 °C) is prepared according to the procedure given in example 8b) (yield: 91 %).

15

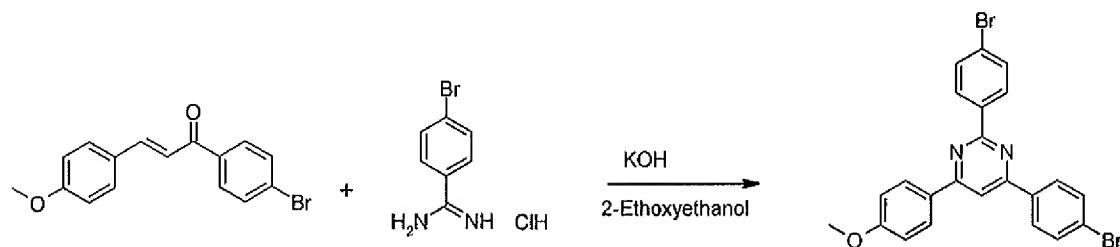


b) The product (melting point: 152-153 °C) is prepared according to the procedure given in example 8b) (yield: 54 %).

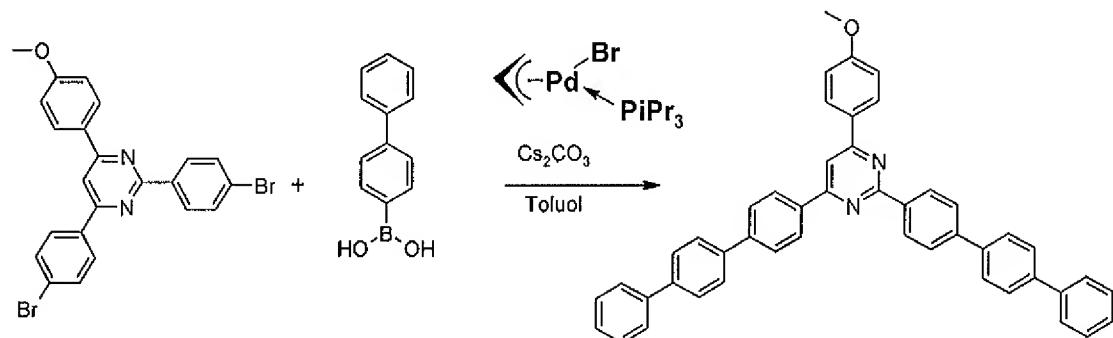


5 c) The product (melting point: 220-221 °C) is prepared according to the procedure given in example 8c) (yield: 80 %).

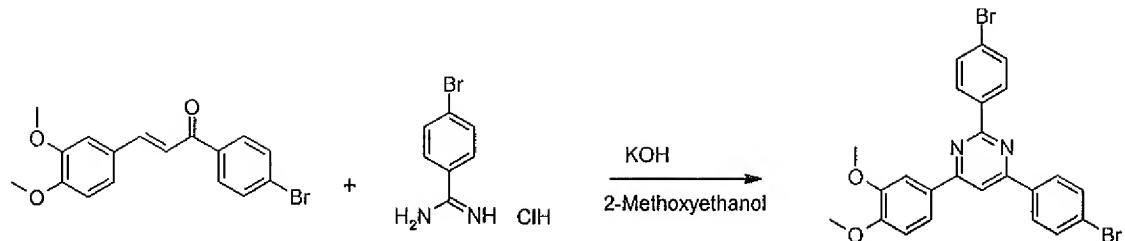
Example 13 (H-6)



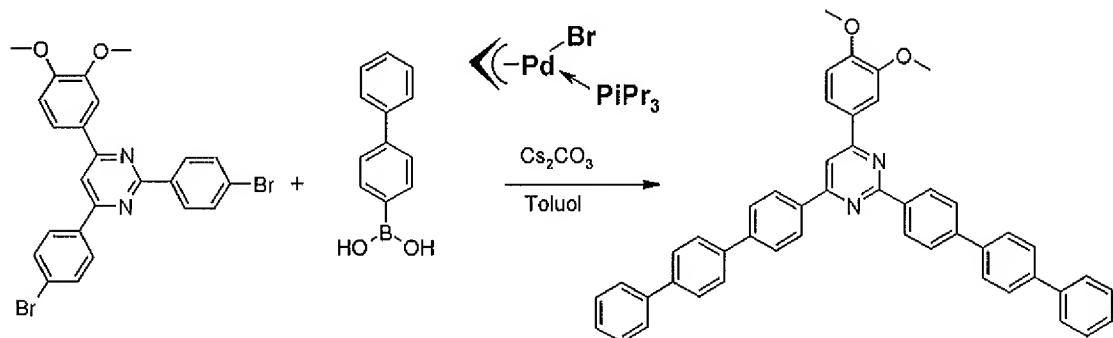
10 a) The product (melting point: 245-246 °C) is prepared according to the procedure given in example 8b) (yield: 42 %).



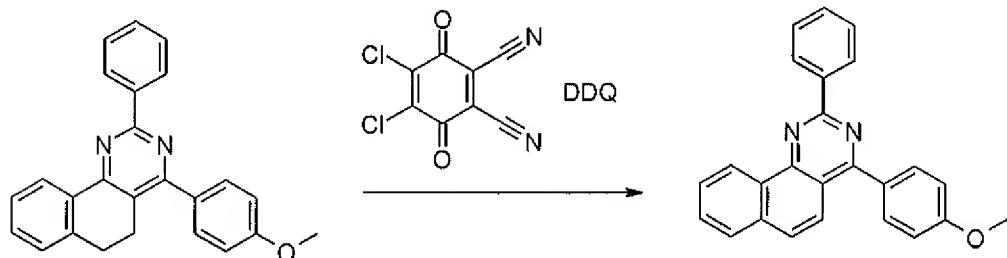
15 b) The product (melting point: 333-334 °C) is prepared according to the procedure given in example 8c) (yield: 75 %).

Example 14 (H-7)

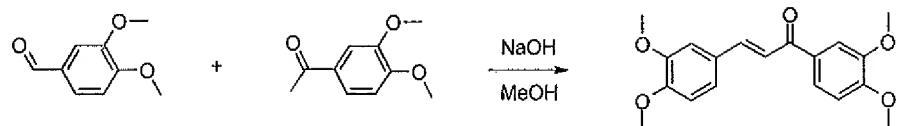
5 a) The product (melting point: 204–205 °C) is prepared according to the procedure given in example 8b) (yield: 46 %).



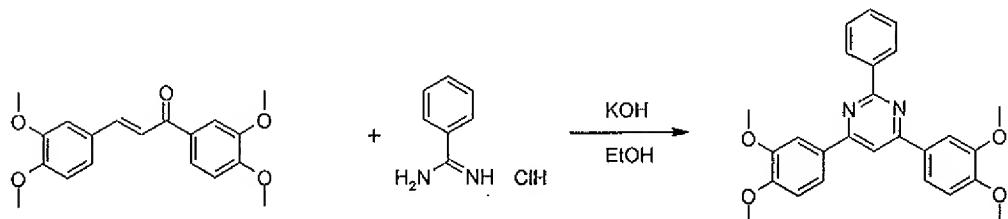
b) The product (melting point: 329–331 °C) is prepared according to the procedure given in example 8c) (yield: 61 %).

10 Example 15

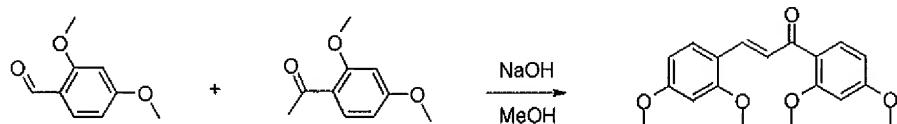
15 To 264 mg (1.0 mmol) of the pyrimidine (see scheme above) in 20 ml chloroform 454 mg (2.0 mmol) DDQ are added. The reaction mixture is refluxed for 1 day. Additional 908 mg (4.0 mmol) of DDQ are added and the reaction mixture is refluxed for additional 2 days. The reaction mixture is filtered on silica gel with dichloromethane. A column chromatography on silica gel with toluene gives the desired product in 61 % yield (mp. 163–165 °C).

Example 16 (J-1)

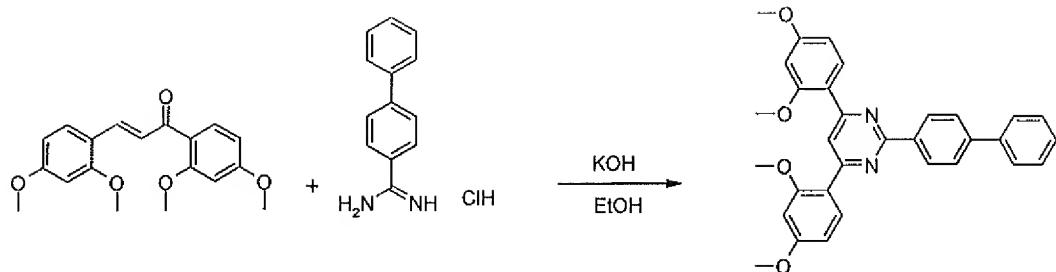
5 a) 10.0 g (57.2 mol) 3,4-dimethoxy-benzaldehyde and 10.3 g (57.2 mol) 3,4-dimethoxy-acetophenone are dissolved under argon in 50 ml methanol. To this solution 0.34 g sodium hydroxide are added. The reaction mixture is stirred for 22 h at 40 °C. The reaction mixture is cooled to 0 °C and the precipitated product is filtered off. The product is dried in vacuum (yield: 18.2 g (97 %), melting point: 108 – 110 °C).



10 b) To 7.00 g (21.3 mmol) of 1,3-bis-(3,4-dimethoxy-phenyl)-2-propen-1-one (1) in 40 ml water free ethanol 1.67 g (10.7 mmol) benzamidine hydrochloride are added under argon. A solution of 1.41 g (25.1 mmol) potassium hydroxide (85%) in 40 ml water free ethanol is added during 20 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 48 h and then poured into water. The product is filtered off and washed with 15 ethanol. A column chromatography on silica gel with toluene gives the desired product J-1 (melting point: 157 – 158 °C).

Example 17 (J-2)

20 a) 10.0 g (57.2 mol) 2,4-dimethoxy-benzaldehyde and 10.3 g 2,4-dimethoxy-acetophenone (57.2 mol) are dissolved under argon in 50 ml methanol. To this solution 0.34 g sodium hydroxide are added. The reaction mixture is stirred for 48 h at 40 °C. The reaction mixture is diluted with water and the precipitated product is filtered off. The product is washed with 25 water and dried in vacuum (melting point: 127 – 129 °C).

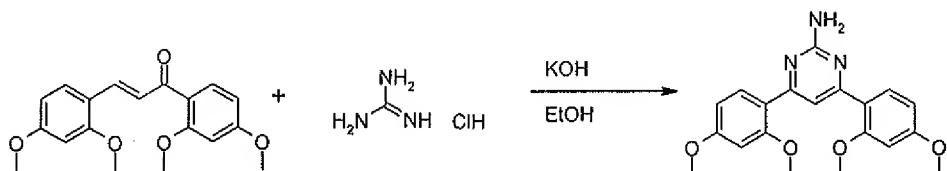


b) To 6.60 g (20.1 mmol) of 1,3-bis-(2,4-dimethoxy-phenyl)-2-propen-1-one (1) in 40 ml water free ethanol 2.34 g (10.1 mmol) biphenyl-benzamidine hydrochloride are added under argon. A solution of 1.41 g (25.1 mmol) potassium hydroxide (85%) in 40 ml water free

5 ethanol is added during 15 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 48 h and then poured into water. The water phase is extracted with dichloromethane. The organic phase is dried with MgSO_4 and the solvent is removed in vacuum. A column chromatography on silica gel with toluene/hexane 3/2 give the desired product J-2 (melting point: 165 – 167 °C).

10

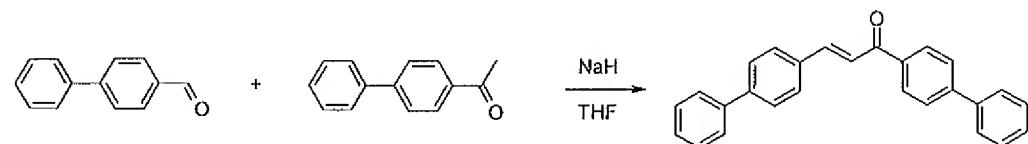
Example 18 (J-4)



To 5.50 g (15.2 mmol) of 1,3-bis-(2,4-dimethoxy-phenyl)-2-propen-1-one (1) in 30 ml water free ethanol 0.87 g (9.14 mmol) guanidine hydrochloride are added under argon. A solution of 1.21 g (21.5 mmol) potassium hydroxide (85%) in 30 ml water free ethanol is added during

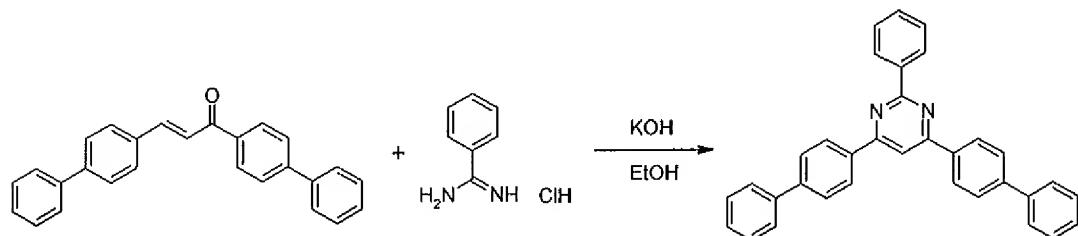
15 15 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 48 h and then poured into water. The water phase is extracted with dichloromethane. The organic phase is dried with MgSO_4 and the solvent is removed in vacuum. A column chromatography on silica gel with dichloromethane/ethyl acetate 2/1 gives the desired product (melting point: 211-213 °C).

Example 19 (J-5)



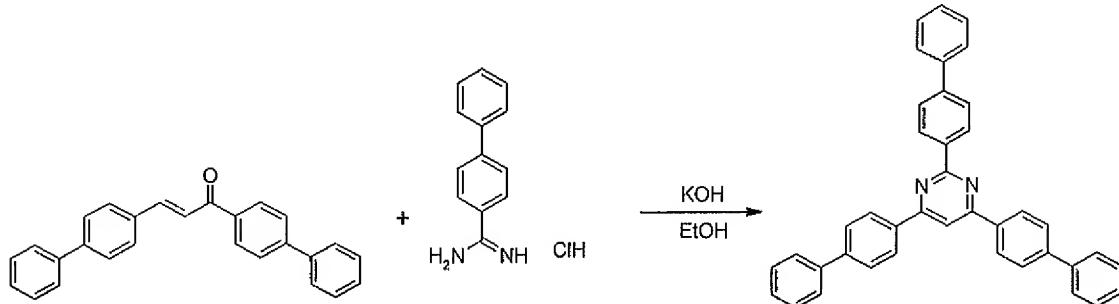
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5 a) To a suspension of 0.69 g sodium hydride in 25 ml terahydrofuran (THF) 5.38 g (27.4 mmol) 4-acetyl-biphenyl are added at 0 °C under argon. After 1 h 5.00 g (27.4 mmol) 4-biphenylcarbadehyde in 25 ml THF are added. The reaction mixture is stirred for 19 h at 25 °C. The formed precipitate is filtered off and washed with THF. The product is refluxed for 1 h in 100 ml iso-propanol and 30 ml water. The product is filtered off and dried in vacuum.



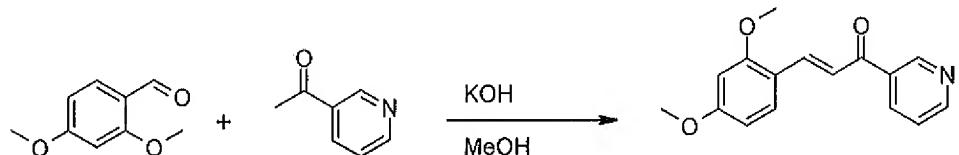
10 b) The desired product J-5 (melting point: 245 – 246°C) is prepared according to the procedure given in example 17b).

Example 20 (J-6)



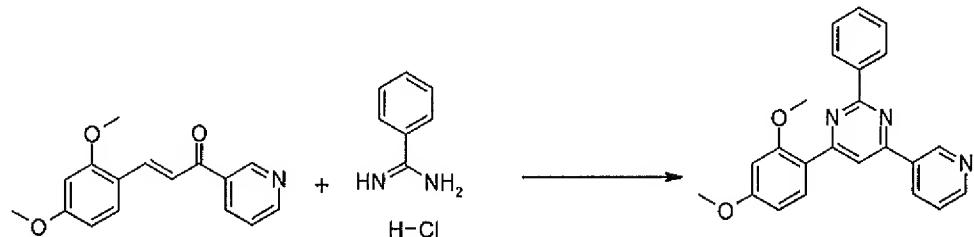
15 The product J-6 (melting point: 243 – 246°C) is prepared according to the procedure given in example 17b).

Example 21 (J-7)



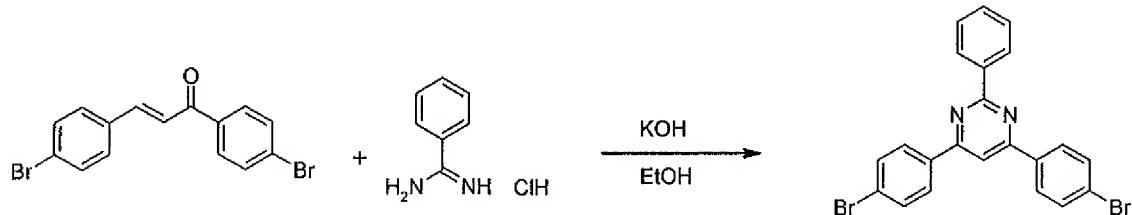
20 a) 18.3 g (0.100 mol) 2,4-dimethoxy-benzaldehyde and 12.1 g (0.100 mol) 3-acetyl pyridine are dissolved under argon in 100 ml methanol. To this solution 0.66 g sodium hydroxide are added. The reaction mixture is refluxed for 18 h. The reaction mixture is poured into water and extracted with dichloromethane. The organic phase is dried with MgSO_4 and the solvent

is removed in vacuum. A column chromatography on silica gel with toluene/ethyl acetate 2/1 give the desired product (yield: 7.3 g (27 %)).

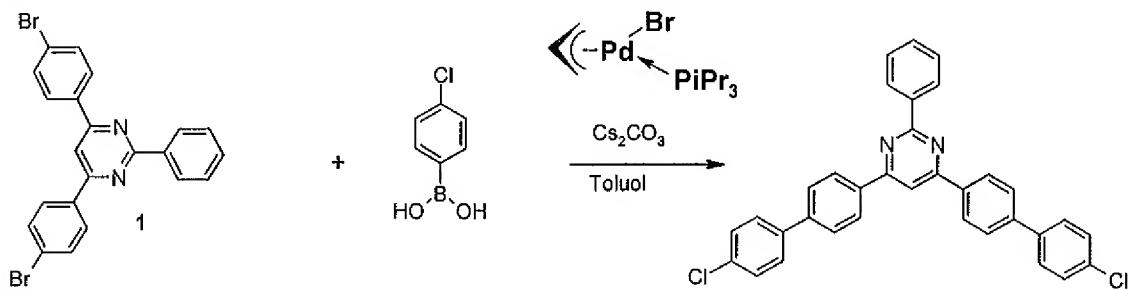


5 b) The product *J*-7 (melting point: 136 °C) is prepared according to the procedure given in example 17b).

Example 22 (K-1)

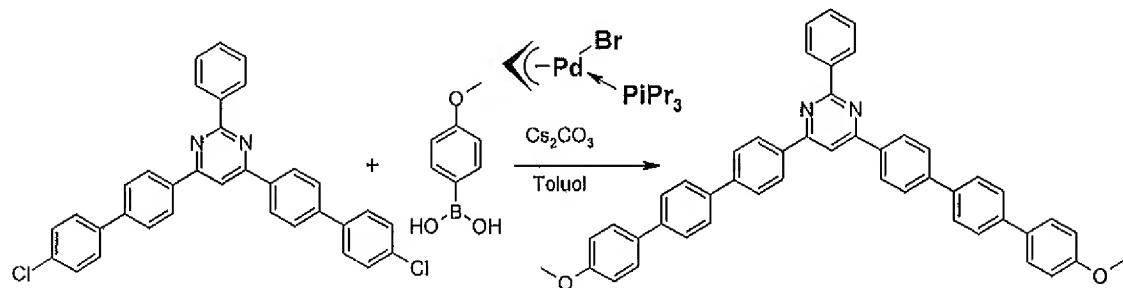


10 a) To 10.0 g (27.3 mmol) of 1,3-bis-(4-bromophenyl)-2-propen-1-one in 70 ml ethanol 2.14 g (13.7 mmol) benzimidine hydrochloride are added under argon. A solution of 1.80 g (32.1 mmol) potassium hydroxide (85%) in 50 ml ethanol is added during 15 min. Dry air is bubbled through the reaction mixture. The reaction mixture is refluxed for 24 h and then is poured into water. The product is filtered off, washed with water and 10 % tartaric acid, crystallized from glacial acetic acid and washed with ethanol (yield: 9.2 g (58 %); melting point: 203 -205 °C).



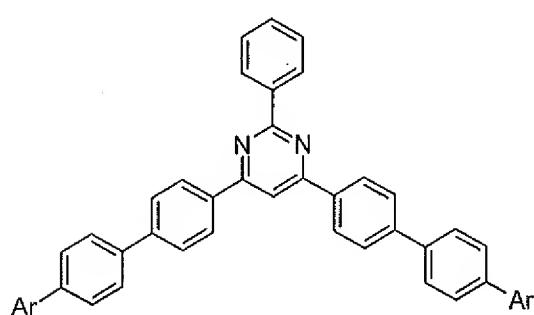
20 b) To 4.45 g (9.55 mmol) 4,6-tris-(p-bromophenyl)-2-phenyl-pyrimidine in 200 ml toluene 2.99 g (19.1 mmol) 4-chloro-phenylboronic acid are added. The suspension is refluxed under argon. Argon is passed through the reaction mixture. 9.33 g (28.6 mmol) of caesium carbonate in 6 ml water are added dropwise to the reaction mixture. Then 0.40 g of the catalyst are added. The reaction mixture is refluxed for 16 h and then the solids are filtered off. The solvent is removed in vacuum. The residue is dissolved in dichloromethane and is

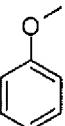
washed with water. The organic phase is dried with magnesium sulfate. The solvent is removed in vacuum. The product is crystallized from 100 ml ethanol (yield: 99 %; melting point: 258 – 259 °C).

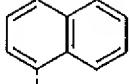
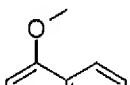
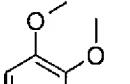
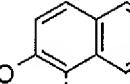
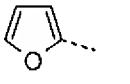
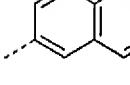
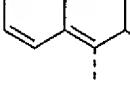
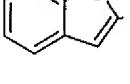
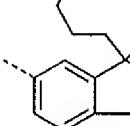
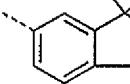
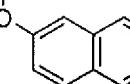


5 c) To 1.00 g (1.89 mmol) 4,6-bis-(p-bromophenyl)-2-phenyl-pyrimidine in 45 ml toluene 287 mg (1.89 mmol) 4-methoxy-biphenylboronic acid are added. Argon is passed through the reaction mixture. 1.23 g (3.78 mmol) of caesium carbonate in 6 ml water is added dropwise to the reaction mixture. Then 20 mg of the catalyst is added. The reaction mixture is refluxed for 18 h. Additional 4-methoxy-biphenylboronic acid and caesium carbonate is added in a 1 to 1 molar ratio until the reaction is complete. The product is filtered off and is washed with ethanol, water and again ethanol. The product is dissolved in boiling DMF, filtered on super gel (Fluka 56678, CAS [91053-39-3]) and crystallized (yield: 0.90 g (71 %); melting point: 345-348 °C).

10 15 In a manner analogous to Example 22 Compounds K-2 to K-15 are obtained.

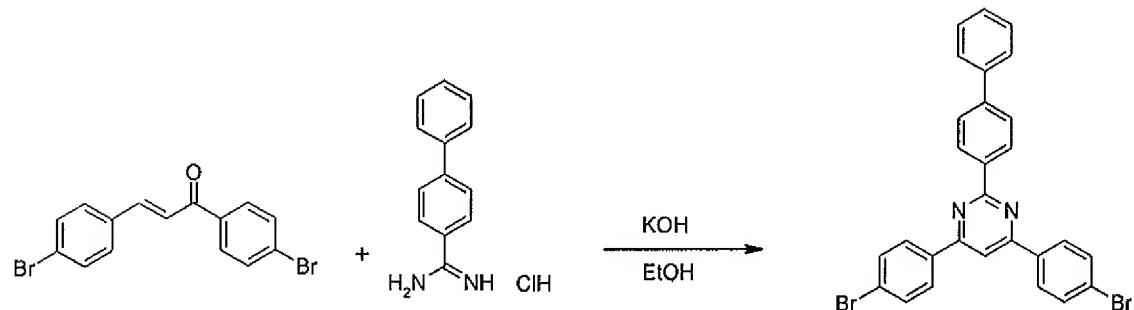


Cpd.	Ar
K-1	 mp. 347 °C

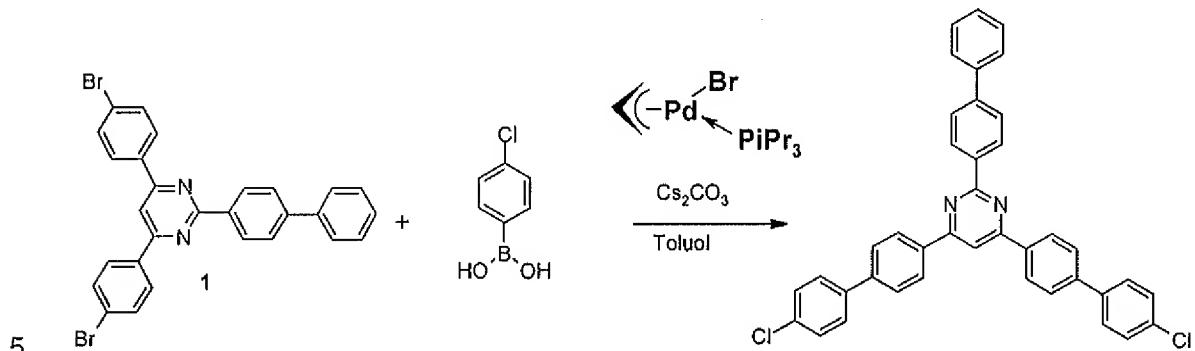
K-2	 mp: 276-279 °C
K-3	
K-4	 mp: 250-251 °C
K-5	
K-6	
K-7	
K-8	
K-9	
K-10	
K-11	
K-12	

K-13	
K-14	
K-15	 mp: 328-330 °C
K-16	

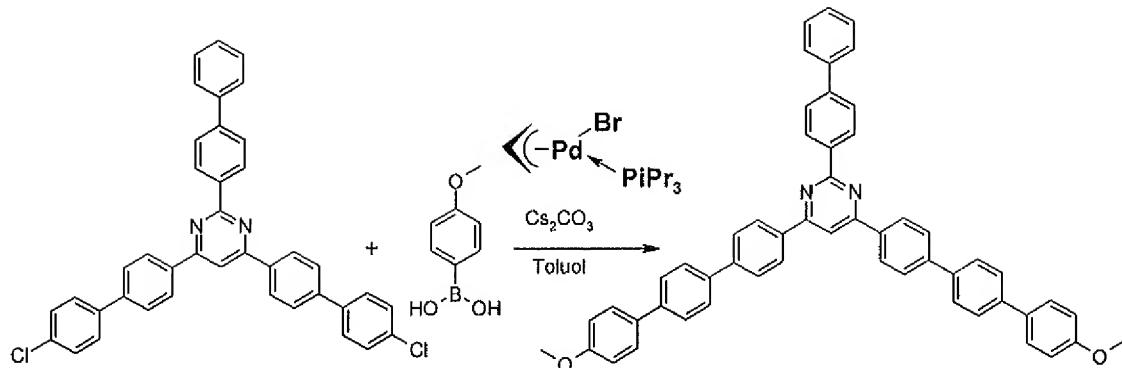
Example 23 (L-1)



a) The product is prepared according to the procedure given in example 22a).

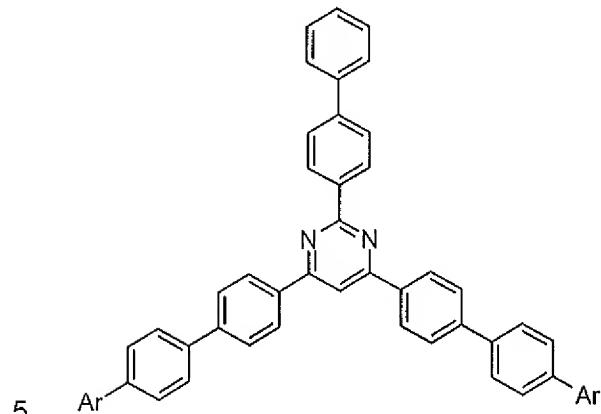


b) The product is prepared according to the procedure given in example 22b).

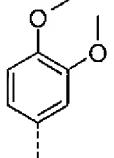
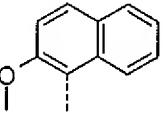
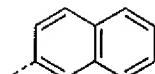
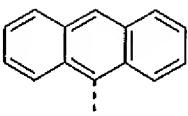
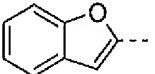
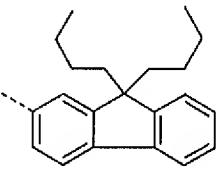
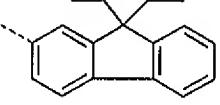
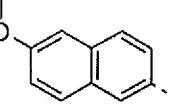
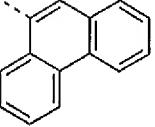
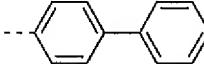


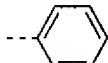
c) The product is prepared according to the procedure given in example 22c).

In a manner analogous to Example 23 Compounds L-2 to L-15 are obtained.



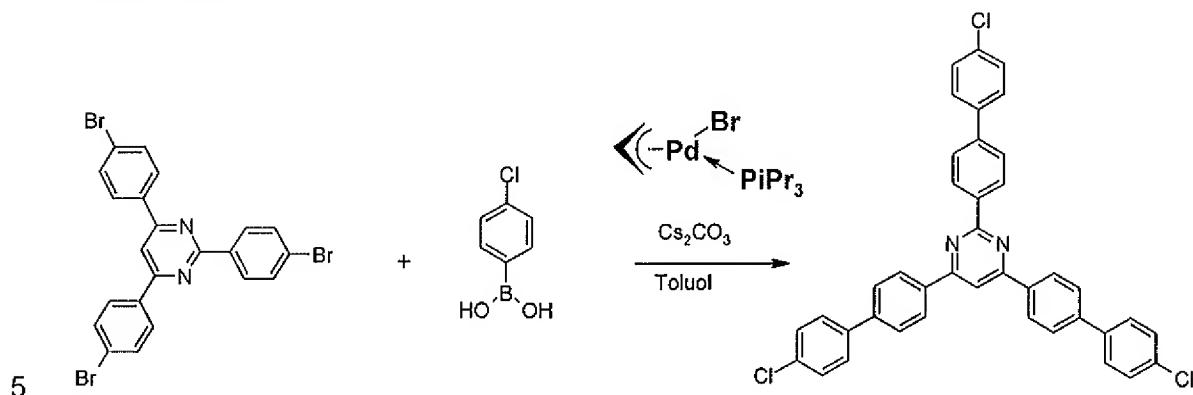
Cpd.	Ar
L-1	
L-2	
L-3	

L-4	
L-5	
L-6	
L-7	
L-8	
L-9	
L-10	
L-11	
L-12	
L-13	
L-14	

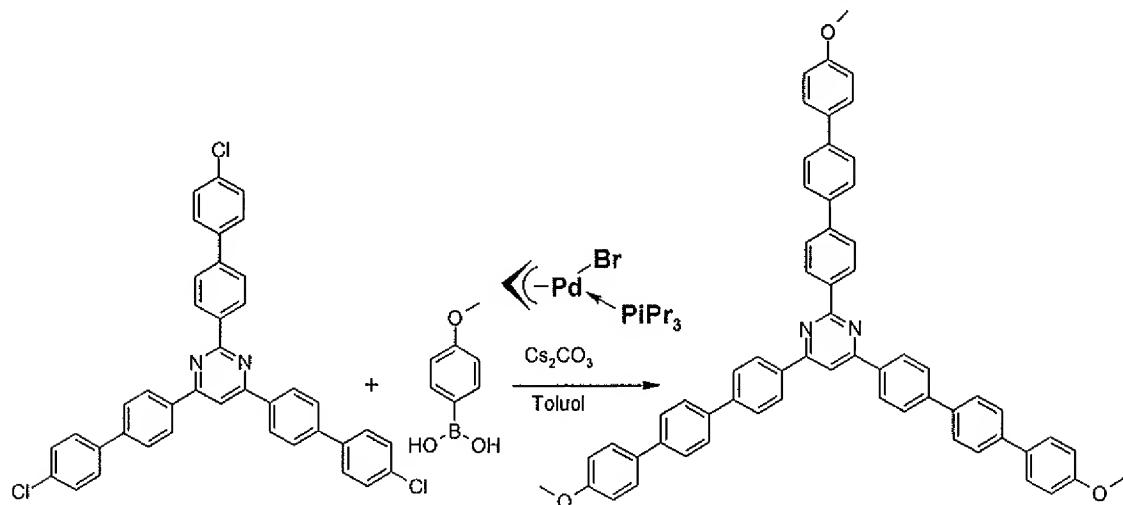
L-15	
L-16 ¹⁾	 mp.: 354 °C

¹⁾ A 100 nm thin film of Compound L-16 showed good film quality and strong photoluminescence at 430 nm (λ_{\max}).

Example 24 (M-1)

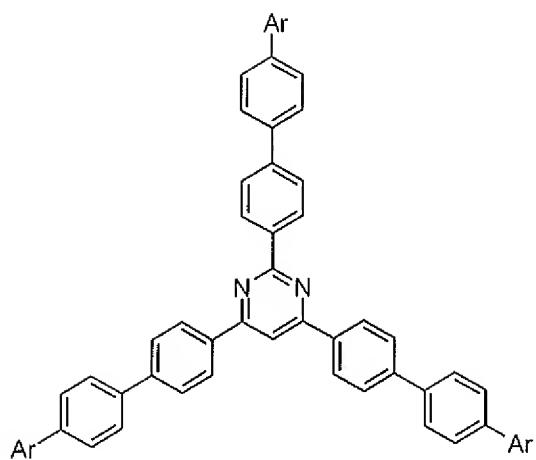


a) The product is prepared according to the procedure given in example 22b).

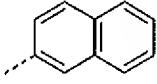
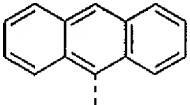
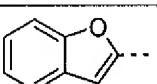
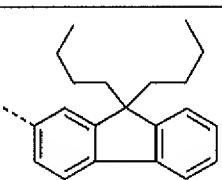
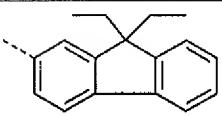
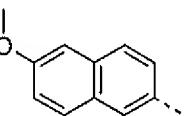
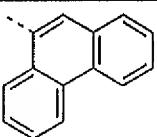
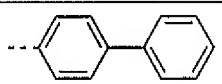
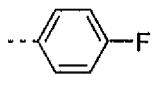
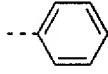


b) The product is prepared according to the procedure given in example 22c).

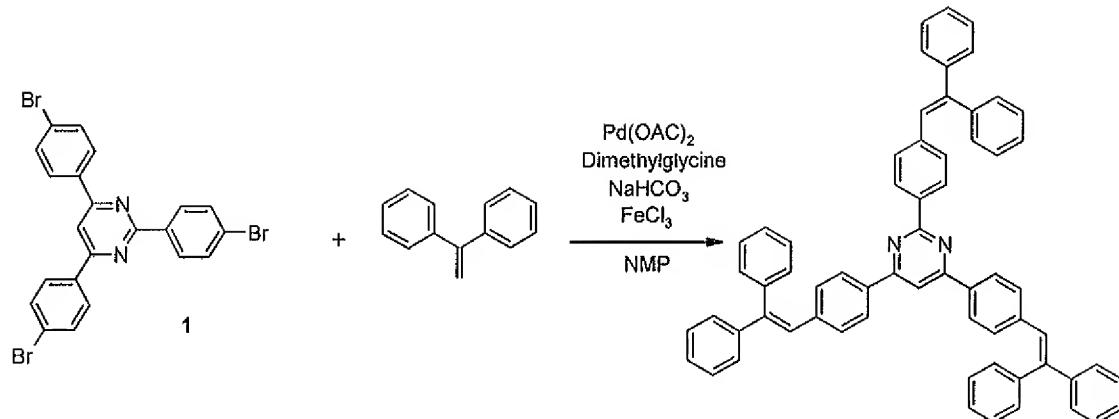
10 In a manner analogous to Example 24 Compounds M-2 to M-15 are obtained.



Cpd.	Ar
M-1	
M-2	
M-3	
M-4	
M-5	
M-6	

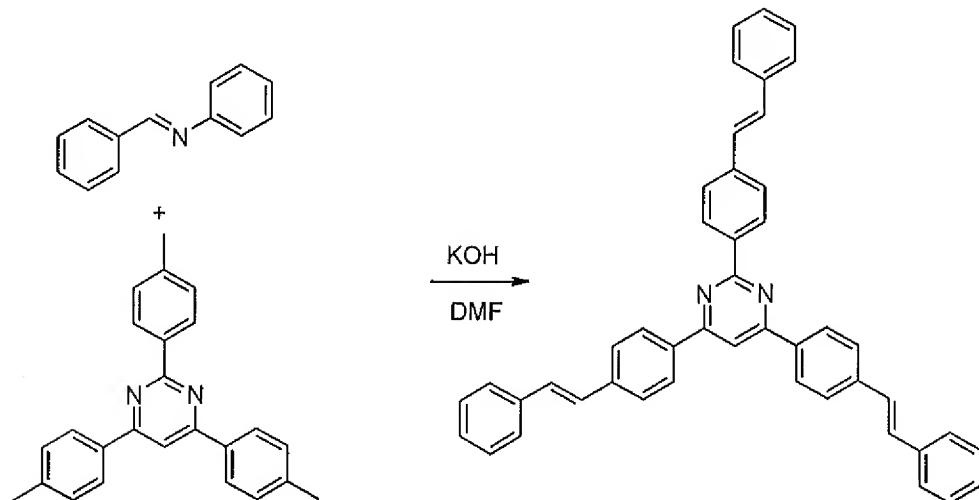
M-7	
M-8	
M-9	
M-10	
M-11	
M-12	
M-13	
M-14	
M-15	
M-16	 mp.: 340 °C

Example 25 (N-1)



1.0 g (1.83 mmol) 2,4,6-tris-(p-bromophenyl)-pyrimidine, 1.49 g (8.26 mmol) 1,1-diphenylethylen, 40 mg Palladium-(II)-acetate, 150 mg (1.47 mmol) dimethylglycine, 1.39 g (16.5 mmol) sodium hydrogen carbonate and 70 mg (0.46 mmol) FeCl_3 are dissolved in 8 ml N-methyl-pyrrolidone. The reaction mixture is heated for 48 h at 150 °C. The reaction mixture is poured into water and 20 % hydrochloric acid is added. The water phase is extracted with dichloromethane. The organic phase is dried with magnesium sulfate. The solvent is distilled off. A column chromatography (silica gel, toluene/hexane 1/3) result in the desired product (119.5 – 120.5 °C).

Example 26



The product is synthesized as described in Example 31 of CH-A-542212.

15

Application Example 1

Present compounds A1, A2, A3, B1, B2, C1, G1 and H1 as light emitting materials,

respectively, 2,5-bis(1-naphthyl)-1,3,4-oxadiazole and a polycarbonate resin in a weight ratio of 5:3:2 are dissolved in tetrahydrofuran, and the solution is spin-coated on a cleaned glass substrate with an ITO electrode to form a light-emitting layer having a thickness of 100 nm. An electrode having a thickness of 150 nm is formed thereon from a magnesium/indium alloy

5 having a magnesium/indium mixing ratio of 10/1, to obtain an organic EL device. The device exhibits light emission with excellent brightness and efficiency at a direct current voltage of 5 V.

Application Example 2

10 Compounds A1, A2, A3, B1, B2, C1, G1 and H1, respectively, are vacuum-deposited on a cleaned glass substrate with an ITO electrode to form a light-emitting layer having a thickness of 100 nm. An electrode having a thickness of 100 nm is formed thereon from a magnesium/silver alloy having a magnesium/silver mixing ratio of 10/1, to obtain an organic EL device. The light-emitting layer is formed by deposition under a vacuum of 10^6 Torr at a

15 substrate temperature of room temperature. The device shows emission having an excellent brightness and efficiency at a direct current voltage of 5 V.

Application Example 3

Compounds A1, A2, A3, B1, B2, C1, G1 and H1, respectively, are dissolved in methylene chloride tetrahydrofuran, and the solution is spin-coated on a cleaned glass substrate with an ITO electrode to form a light-emitting layer having a thickness of 50 nm. Then, aluminum bis(2-methyl-8-quinolinate)(2-naphtolate) is vacuum-deposited to form an electron-injecting layer having a thickness of 10 nm, and an electrode having a thickness of 100 nm is formed thereon from a magnesium/aluminum alloy having a magnesium/aluminum mixing ratio of

25 10/1, to obtain an organic EL device. The light-emitting layer and the electron-injecting layer are formed by deposition under a vacuum of 10^6 Torr at a substrate temperature of room temperature. The device shows an emission having an excellent brightness and efficiency at a direct current voltage of 5 V.

30 **Application Example 4**

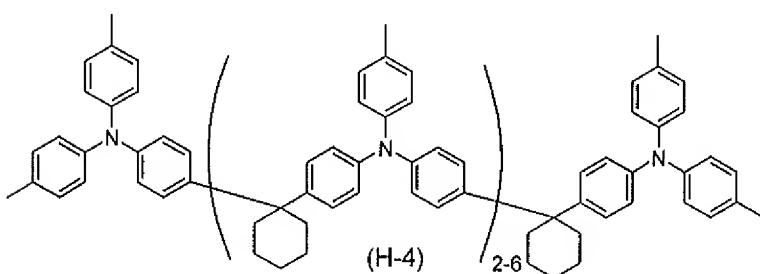
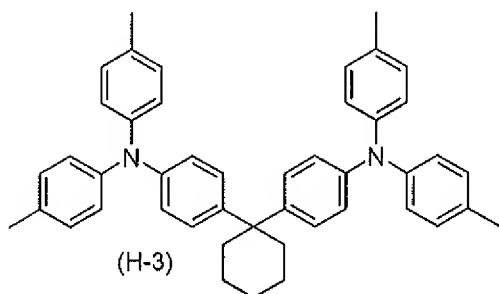
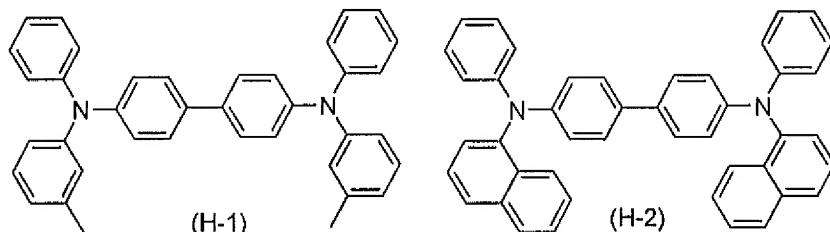
Compounds A1, A2, A3, B1, B2, C1, G1 and H1, respectively, are vacuum-deposited on a cleaned glass substrate with an ITO electrode to form a light-emitting layer having a thickness of 50 nm. Then, aluminum tris(8-hydroxyquinolinate) is vacuum-deposited to form an electron-injecting layer having a thickness of 10 nm and an electrode having a thickness

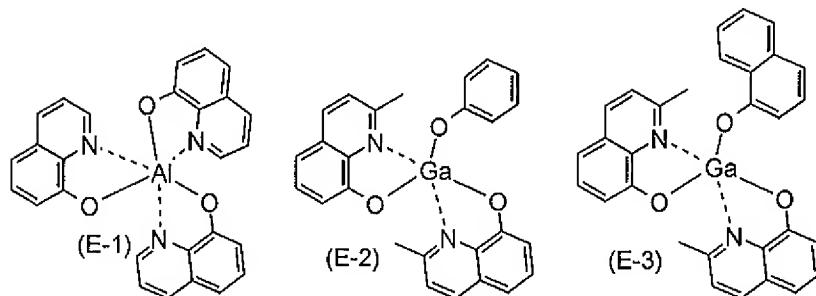
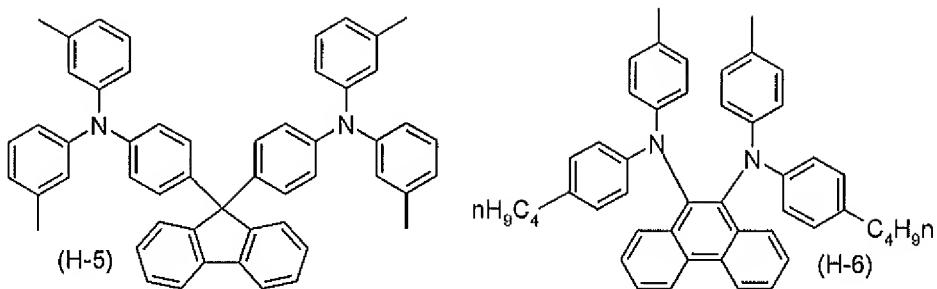
35 of 100 nm is formed thereon from an aluminum/lithium alloy having an aluminum/lithium mixing ratio of 50/1, to obtain an organic EL device. A hole-injecting layer and the light-

emitting layer are formed by deposition under a vacuum of 10^6 Torr at a substrate temperature of room temperature. The device shows a light emission having an excellent brightness and efficiency at a direct current voltage of 5 V.

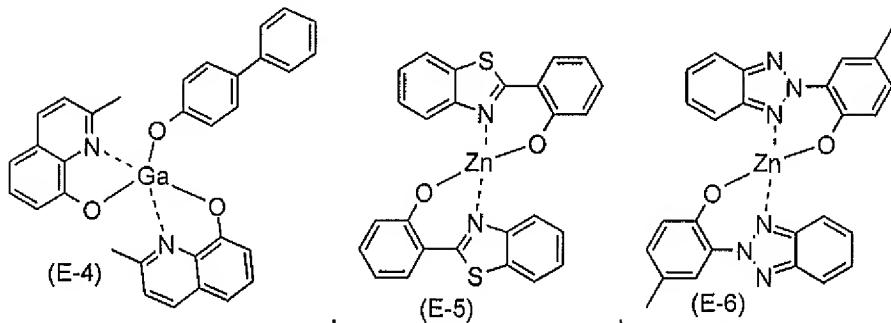
5 **Application Example 5**

One of hole-injecting materials (H-1) to (H-6) is vacuum-deposited on a cleaned glass substrate with an ITO electrode, to form a hole-injecting layer having a thickness of 30 nm. Then, one of light-emitting materials A1, A2, A3, B1, B2, C1, G1 and H1, respectively is vacuum-deposited to form a light-emitting layer having a thickness of 30 nm. Further, one of 10 electron-injecting materials (E-1) to (E-6) is vacuum-deposited to form an electron-injecting layer having a thickness of 30 nm. An electrode having a thickness of 150 nm is formed thereon from a magnesium/silver alloy having a magnesium/silver mixing ratio of 10/1, to obtain an organic EL device. Each layer is formed under a vacuum of 10^6 Torr at a substrate temperature of room temperature. All the organic EL devices obtained in these Examples 15 shows high brightness and efficiency.





5



Application Example 6

On a cleaned glass substrate with an ITO electrode, 4,4',4"-tris(N-(3-methylphenyl)-N-phenylamino)triphenylamine is vacuum-deposited to form a first hole-injecting layer having a thickness of 25 nm. Further, a hole-injecting material (H-1) is vacuum-deposited to form a second hole-injecting layer having a thickness of 5 nm. Then, compounds A1, A2, A3, B1, B2, C1, G1 and H1, respectively, as light-emitting materials are vacuum-deposited to form a light-emitting layer having a thickness of 20 nm. Further, an electron-injecting material (E-1) is vacuum-deposited to form an electron-injecting layer having a thickness of 30 nm. Then, an electrode having a thickness of 150 nm is formed thereon from a magnesium/silver alloy having an magnesium/silver mixing ratio of 10/1, to obtain an organic EL device. The device shows emission having an outstanding brightness and efficiency at a direct current voltage of 5 V.

Application Example 7

On a cleaned glass substrate with an ITO electrode, 4,4',4''-tris(N-(1-naphthyl)-N-phenylamino)triphenylamine is vacuum-deposited to form a first hole-injecting layer having a thickness of 25 nm. Further, a hole-injecting material (H-2) is vacuum-deposited to form a second hole-injecting layer having a thickness of 5 nm. Then, compounds A1, A2, A3, B1, B2, C1, G1 and H1, respectively, as light-emitting materials are vacuum-deposited to form a light-emitting layer having a thickness of 20 nm. Further, an electron-injecting material (E-5) is vacuum-deposited to form an electron-injecting layer having a thickness of 30 nm. Then, an electrode having a thickness of 150 nm is formed thereon from a magnesium/silver alloy having an magnesium/silver mixing ratio of 10/1, to obtain an organic EL device. The device shows an emission having a outstanding brightness and efficiency at a direct current voltage of 5 V.

Application Example 8

A hole-injecting material (H-5) is vacuum-deposited on a cleaned glass substrate with an ITO electrode to form a hole-injecting layer having a thickness of 20 nm. Then, compounds A1,

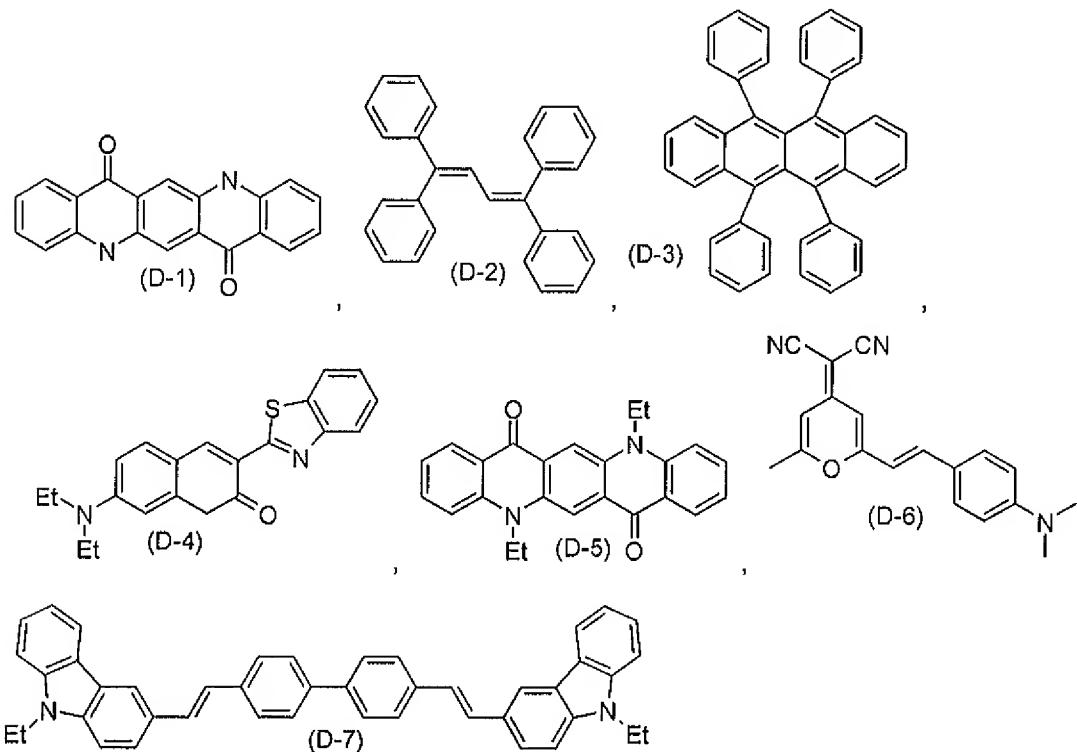
5 A2, A3, B1, B2, C1, G1 and H1, respectively, as light-emitting materials are vacuum-deposited to form a light-emitting layer having a thickness of 20 nm. Further, an electron-injecting material (E-2) is vacuum-deposited to form a first electron-injecting layer having a thickness of 20 nm. Then, an electron-injecting material (E-5) is vacuum-deposited to form a second electron-injecting layer having a thickness of 10 nm, and an electrode having a

10 thickness of 150 nm is formed thereon from a magnesium/silver alloy having an magnesium/silver mixing ratio of 10/1, to obtain an organic EL device. The device shows light emission having an excellent brightness and efficiency at a direct current voltage of 5 V.

Application Example 9

15 An organic EL device is prepared in the same manner as in Example 5 except that the light-emitting layer is replaced with a 30 nm thick light-emitting layer formed by vacuum-depositing compounds A1, A2, A3, B1, B2, C1, G1 and H1, respectively, and one of the dopant compounds (D-1) to (D-7) in a weight ratio of 100:1. All the organic EL devices obtained in these Examples shows high brightness characteristics and gives intended light emission

20 colors.



Application Example 10

An organic EL device is prepared in the same manner as in Example 5 except that the light-emitting layer is replaced with a 30 nm thick light-emitting layer formed by vacuum-depositing compounds A1, A2, A3, B1, B2, C1, G1 and H1, respectively and one of Compounds (D-1) to (D-7) in a weight ratio of 100:1. All the organic EL devices obtained in these Examples shows high brightness characteristics, or a maximum brightness and gives intended light emission colors.

10 Application Example 11

A hole-injecting material (H-2) is vacuum-deposited on a cleaned glass substrate with an ITO electrode to form a hole-injecting layer having a thickness of 30 nm. Then, 4,4'-bis(α,α -diphenylvinyl)biphenyl and a light-emitting material selected from compounds A1, A2, A3, B1, B2, C1, G1 and H1 for a light-emitting layer are vacuum-deposited in a weight ratio of 100:5 to form a light-emitting layer having a thickness of 30 nm. Further, an electron-injecting material (E-3) is vacuum-deposited to form an electron-injecting layer having a thickness of 30 nm. Then, an electrode having a thickness of 150 nm is formed thereon from a magnesium/silver alloy having an magnesium/silver mixing ratio of 10/1, to obtain an organic EL device. The device shows an emission having an outstanding brightness and efficiency at a direct current voltage of 5 V.

Application Example 12

An organic EL device is prepared in the same manner as in Example 11 except that the light-emitting layer is replaced with a 30 nm thick light-emitting layer formed by vacuum-depositing 5 aluminum tris(8-hydroxyquinolinate) and one of the light-emitting materials A1, A2, A3, B1, B2, C1, G1 and H1 in a weight ratio of 100:3. All the organic EL devices obtained in these Examples shows high brightness characteristics at a direct current of 5 V.

The organic EL devices obtained in the Application Examples of the present invention show 10 an excellent light emission brightness and achieved a high light emission efficiency. When the organic EL devices obtained in the above Examples are allowed to continuously emit light at 3 (mA/cm²), all the organic EL devices remain stable. Since the light-emitting materials of the present invention have a very high fluorescence quantum efficiency, the organic EL devices using the light-emitting materials achieved light emission with a high brightness in a 15 low electric current applied region, and when the light-emitting layer additionally uses a doping material, the organic EL devices are improved in maximum light emission brightness and maximum light emission efficiency. Further, by adding a doping material having a different fluorescent color to the light-emitting material of the present invention, there are obtained light-emitting devices having a different light emission color. The organic EL devices 20 of the present invention accomplish improvements in light emission efficiency and light emission brightness and a longer device life, and does not impose any limitations on a light-emitting material, a dopant, a hole-injecting material, an electron-injecting material, a sensitizer, a resin and an electrode material used in combination and the method of producing the device. The organic EL device using the material of the present invention as a 25 light-emitting material achieves light emission having a high brightness with a high light emission efficiency and a longer life as compared with conventional devices. According to the light-emitting material of the present invention and the organic EL device of the present invention, there can be achieved an organic EL device having a high brightness, a high light emission efficiency and a long life.